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[Handwritten signature]

Schnizer, Holly
Wednesday, January 02, 2002 1:21 PM
STIC-Biotech/ChemLib
seq. search request for appl. no. 09/444,281

Please search the commercial and interference databases for

SEQ ID NOs: 27, 35, and 36 and polynucleotides encoding SEQ ID NOs: 27, 35, and 36

Thank you.

Holly Schnizer
AU 1653
CM1-10B05
305-3722
mailbox: CM1-9B01

CRPG

if Contact:
Searcher: Sheppard
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Searcher Prep/Review: _____
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TYPE OF SEARCH:
NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____
Litigation: _____
Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST(where applic.)
STN: _____
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Questel/Orbit: _____
DRLink: _____
Lexis/Nexis: _____
Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____

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PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
PS Claim 11; Page 88; 129pp; English.
XX
CC AAY24549 to AAY24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): RX₁XX₂XB (I), BX₁XX₂XB
CC (II), BBX₁XX₂XB (III), BX₁XX₂BBB(AA)nMBB(AA) (IV), BX₁XX₂BBB(AA)nM
CC (V), LBnX₁XX₂BBB(AA)nX₃RR (VI), LKX₁XX₂BBB(AA)nX₃RR (VII), BBX₁XX₂BBB(AA)nM
CC (VIII), where 2 = P or V; X = hydrophobic residue, preferably W, B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC derived from the analogues may be used similarly; the compounds may
CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.
XX
SQ Sequence 12 AA;
XX

Query Match 100.0%; Score 86; DB 19; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
| | | | | | | | | |
Db 1 ILRPMWPMWRK 12

RESULT 2
AAY94496
ID AAY94496 standard; Peptide; 12 AA.
XX
AC AAY94496;
XX
DT 20-SEP-2000 (first entry)
XX
DE MBI-11B7 peptide derived from indolicidin.
XX
KM Cellulose binding domain; CBD; cationic peptide;
KM MBI-11B7; indolicidin; bovine.
XX
OS Bos taurus.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PE 19-NOV-1999; 99WO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld P.
XX
DR WPI: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
XX
PS Disclosure; Page 24; 73pp; English.

XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is the MBI-11B7 peptide. MBI-11B7 is a cationic peptide derived
CC from modifications of indolicidin.
XX
SQ Sequence 12 AA;
XX

Query Match 100.0%; Score 86; DB 21; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
| | | | | | | | | |
Db 1 ILRPMWPMWRK 12

RESULT 3
AAY91791
ID AAY91791 standard; Peptide; 12 AA.
XX
AC AAY91791;
XX
DT 06-JUN-2000 (first entry)
XX
DE Amino acid sequence of cationic peptide MBI 11B7CN.
XX
KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KM leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KM multidrug resistance.
XX
OS Synthetic.
XX
PN WO9965506-A2.
XX
PD 23-DEC-1999.
XX
PE 14-JUN-1999; 99WO-CA00552.
XX
PR 12-JUN-1998; 98US-0096541.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Friedland HD, Krieger TJ, Taylor R, Erle D, Fraser JR, West MHP.
XX
DR WPI: 2000-223549/19.
XX
PT Novel pharmaceutical composition containing optionally activated
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
XX
PS Claim 1; Page 14; 94pp; English.
XX
CC This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxyalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.
XX
SQ Sequence 12 AA;

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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:26 ; Search time 53.46 Seconds
(without alignments)
16.627 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPMWRRK 12

Scoring table: BIOSUM62
Gapop 10.0, Gapext 0.5

Searched: 522463 seqs, 74073290 residues
Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 08
Maximum Match 1008
Listing first 45 summaries

Database :

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21: /SIDSR/gcgdata/geneseq/geneseq/AA2000.DAT.*
22: /SIDSR/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	86	100.0	12	19	AAV24550
2	86	100.0	12	21	AAV94496
3	86	100.0	12	21	AAV91791
4	86	100.0	20	19	AAV24553
5	86	100.0	20	21	AAV91797
6	86	100.0	21	19	AAV24552
7	86	100.0	21	19	AAV24554
8	86	100.0	21	19	AAV66376
9	86	100.0	21	21	AAV91796
10	86	100.0	21	21	AAV91798
11	86	100.0	27	19	AAV66363

12	86	100.0	28	21	AAV91800
13	83	96.5	12	19	AAV24567
14	83	96.5	12	21	AAV91788
15	82	95.3	12	19	AAV24594
16	82	95.3	12	19	AAV66364
17	82	95.3	12	21	AAV91817
18	82	95.3	12	21	AAV91841
19	81	94.2	12	19	AAV24605
20	81	94.2	12	19	AAV24595
21	81	94.2	12	21	AAV91842
22	81	94.2	12	21	AAV91852
23	80	93.0	12	19	AAV24596
24	80	93.0	12	19	AAV24603
25	80	93.0	12	19	AAV24604
26	80	93.0	12	21	AAV91843
27	80	93.0	12	21	AAV91850
28	80	93.0	12	21	AAV91851
29	78	90.7	12	19	AAV24598
30	78	90.7	12	19	AAV24601
31	78	90.7	12	19	AAV66361
32	78	90.7	12	21	AAV91785
33	78	90.7	12	21	AAV91845
34	78	90.7	12	21	AAV91848
35	78	90.7	13	19	AAV24565
36	78	90.7	13	21	AAV24586
37	77	89.5	12	19	AAV24586
38	77	89.5	12	21	AAV91786
39	75	87.2	11	19	AAV24569
40	75	87.2	11	21	AAV91790
41	75	87.2	12	19	AAV24580
42	75	87.2	12	21	AAV91804
43	75	87.2	13	18	AAV12873
44	75	87.2	13	18	AAV12895
45	75	87.2	13	18	AAV12896

ALIGNMENTS

RESULT 1
ID AAV24550 standard; peptide; 12 AA.
AC AAV24550;
XX
DT 18-AUG-1999 (first entry)
XX
DE Indolicidin analogue #2.
XX
KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.
XX
OS Synthetic.
XX
PN WO9807745-A2.
XX
PD 26-FEB-1998.
XX
PF 21-AUG-1997; 97WO-US14779.
XX
PR 13-JAN-1997; 97US-0034949.
XX
PR 21-AUG-1996; 96US-0024754.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Effle D, Fraser JR, Krieger TU, Taylor R, West MH;
XX
DR WPI; 1998-169090/15.
XX
PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also

Query Match 100.0%; Score 86; DB 21; Length 12;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRK 12
DB 1 ILRWPMPWRK 12

RESULT 4
AAV24553
ID AAV24553 standard; peptide: 20 AA.
XX AAV24553;
AC AAV24553;
XX 18-AUG-1999 (first entry)
DT 18-AUG-1999 (first entry)
XX Indolicidin analogue #5.
DE Indolicidin; bacterial infection; photo-oxidised solubiliser;
XX antimicrobial; antibiotic; antitarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KM food; technical material.
XX Synthetic.
OS Synthetic.
PN WO9807745-A2.
XX 26-FEB-1998.
PD 26-FEB-1998.
XX 21-AUG-1997; 97WO-US14779.
PF 21-AUG-1997; 97WO-US14779.
XX 13-JAN-1997; 97US-0034949.
PR 21-AUG-1996; 96US-0024754.
XX (MICR-) MICROLOGIX BIOTECH INC.
PA (MICR-) MICROLOGIX BIOTECH INC.
XX Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;
PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;
DR WPI: 1998-169090/15.
XX New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX Claim 11; Page 88; 129pp; English.

AAV24549 to AAV24615 represent indolicidin analogues of formulae
(I)-(VIII) containing up to 25 amino acids (aa): RZXZXZXB (I), BXZXZXZB
(II), BBBZXZXZXB (III), BXZXZXZBBn(A)nmlBBAGS (IV), BXZXZXZBB(A)nM
(V), LBBnZXZXZnR (VI), LKnZXZXZXRK (VII) and BBZXZXZBBB (VIII).
Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
in (VIII) at least 2 X = F or Y. The analogues are used to treat
infections caused by bacteria (Gram positive or negative, or anaerobic);
fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
trematodes) or viruses. Typical of very many pathogens that can be
controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
derived from the analogues may be used similarly; the compounds may
also be prepared from antibiotics or antitarrhythmic agents. The analogues
may be used therapeutically or to coat medical devices; also they are
useful as surface disinfectants, as additives to shampoo or soaps, as
insecticides or herbicides, or as preservatives for foods and technical
materials. The analogues are administered by injection, lavage, orally
or topically, generally at 0.1-50 mg/Kg. These analogues have a broader
spectrum of activity than indolicidin and modification as compounds
reduces their toxicity.

SQ Sequence 20 AA;

Query Match 100.0%; Score 86; DB 19; Length 20;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRK 12
DB 1 ILRWPMPWRK 12

RESULT 5
AAV91797
ID AAV91797 standard; peptide: 20 AA.
XX AAV91797;
AC AAV91797;
XX 06-JUN-2000 (first entry)
DT 06-JUN-2000 (first entry)
XX Amino acid sequence of cationic peptide MBI 11B17CN.
DE Amino acid sequence of cationic peptide MBI 11B17CN.
XX Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KM multidrug resistance.
XX Synthetic.
OS Synthetic.
PN WO9965506-A2.
XX 23-DEC-1999.
PD 23-DEC-1999.
XX 14-JUN-1999; 99WO-CA00552.
PF 14-JUN-1999; 99WO-CA00552.
XX 12-JUN-1998; 98US-0096541.
PR 12-JUN-1998; 98US-0096541.
XX (MICR-) MICROLOGIX BIOTECH INC.
PA (MICR-) MICROLOGIX BIOTECH INC.
XX Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
DR WPI: 2000-223549/19.
XX Novel pharmaceutical composition containing optionally activated
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
XX Disclosure: Page 15; 94pp; English.

This sequence represents a cationic peptide amino acid sequence, which
can be used in the pharmaceutical composition of the invention. The
invention relates to a pharmaceutical composition containing at least one
activated polyoxyalkylene (APO)-modified cationic peptide. The
modification of peptides with APO increases their activity against tumour
cells, including those with a multidrug resistant phenotype. The
pharmaceutical composition can be used to treat tumours, specifically
lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
cervix, uterus, skin, prostate, liver and colon.

SQ Sequence 20 AA;

Query Match 100.0%; Score 86; DB 21; Length 20;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRK 12
DB 1 ILRWPMPWRK 12

RESULT 6
AAV24552

ID AAY24552 standard; peptide; 21 AA.
 AC AAY24552;
 XX
 DT 18-AUG-1999 (first entry)
 XX
 DE Indolicidin analogue #4.
 XX
 KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KW antimicrobial; antibiotic; antitarrhythmic; surface disinfectant;
 KW additive; Shampoo; soap; insecticide; herbicide; preservative;
 KW food; technical material.
 OS Synthetic.
 XX
 PN MO9807745-A2.
 XX
 PD 26-FEB-1998.
 XX
 PF 21-AUG-1997; 97WO-US14779.
 XX
 PR 13-JAN-1997; 97US-0034949.
 PR 21-AUG-1996; 96US-0024754.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Ernie D, Fraser JR, Krieger TU, Taylor R, West MH;
 PI
 DR WPI: 1998-169090/15.
 XX
 PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid - vector, transformed cells and antibodies, also
 PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.
 XX
 PS Claim 11; Page 88; 129pp; English.
 XX
 AA AAY24549 to AAY24615 represent indolicidin analogues of formulae
 (I)-(VIII) containing up to 25 amino acids (aa): R₁XX₁XB (I), B₁XX₁XB
 (II), B₁B₂XX₁XB (III), B₁B₂XX₁B₃BB₁(AA)N₁LB₁B₂AGS (IV), B₁XX₁XB₁(AA)₁N
 (V), LB₁N₁XX₁XB₁NR₁ (VI), LK₁XX₁XB₁RR₁ (VII) and B₁XX₁XB₁BB₁ (VIII).
 Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
 preferably R or K; AA = any aa; n = 0 or 1; In (II), at least 1 Z = V;
 in (VIII) at least 2 X = F or Y. The analogues are used to treat
 infections caused by bacteria (gram positive or negative, or anaerobic);
 fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 trématodes) or viruses. Typical of very many pathogens that can be
 controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
 derived from the analogues may be used similarly; the compounds may
 also be prepared from the analogues or antitarrhythmic agents. The compounds
 may be used therapeutically or to coat medical devices; also they are
 useful as surface disinfectants, as additives to shampoo or soaps, as
 insecticides or herbicides, or as preservatives for foods and technical
 materials. The analogues are administered by injection, lavage, orally
 or topically, generally at 0.1-50 mg/Kg. These analogues have a broader
 spectrum of activity than indolicidin and modification as compounds
 reduces their toxicity.
 XX
 Sequence 21 AA;
 50

ID AY24554 standard; peptide; 21 AA.
AC AMY24554;
DT 18-AUG-1999 (first entry)
DE Indolicidin analogue #6.
KW Indolicidin; bacterial infection; photo-oxidised solibiliser;
KW antimicrobial; antibiotic; antitarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.
OS Synthetic.
PN WC0907745-A2.
PD 26-FEB-1998.
PE 21-AUG-1997; 97WO-US14779.
PX 13-JAN-1997; 97US-0034949.
PR 21-AUG-1996; 96US-0024754.
PA (MICR-) MICROLOGIX BIOTECH INC.
PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;
DR WPI: 1998-169090/15.
PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxalylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX Claim 11; Page 88; 129pp; English.

AY24549 to AY24615 represent indolicidin analogues of formulae
(I)-(VIII) containing up to 25 amino acids (aa): RX2XXXB (I), BX2XZXB
(II), BBK2XXXZB (III), BX2XZXB(BN)(AA)nMILBAGS (IV), BX2XZXB(AA)nM
(V), LBBAZX2XZXRK (VI), LKN2XZXRKRK (VII) and BRX2XZXB(B (VIII)).
Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
preferably R or K; AA = any aa; n = 0 or 1; in (II) at least 1 Z = V;
in (VIII) at least 2 X = F or Y. The analogues are used to treat
infections caused by bacteria (Gram positive or negative, or anaerobic);
fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
trematodes) or viruses. Typical of very many pathogens that can be
controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
derived from the analogues may be used similarly: the compounds may
also be prepared from antibiotics or antiarrhythmic agents. The analogues
may be used therapeutically or to coat medical devices; also they are
useful as surfactant disinfectants, as additives to shampoo or soaps, as
insecticides or herbicides, or as preservatives for foods and technical
materials. The analogues are administered by injection, lavage, orally
or topically, generally at 0.1-50 mg/kg. These analogues have a broader
spectrum of activity than indolicidin and modification as compounds
reduces their toxicity.

Sequence 21 AA;

RESULT 8
 ID AAM6376 standard; peptide: 21 AA.
 AC AAM6376;
 DT 12-JAN-1999 (first entry)
 DE Cationic peptide of claim 15 #3.
 KM Indolicidin analogue; resistance; cationic peptide; antibiotic;
 KM bacterial infection; tolerance; antibacterial; microorganism;
 KM bacteria; fungus; parasite; virus.
 OS Synthetic.
 PN WO9840401-A2.
 PD 17-SEP-1998.
 PF 10-MAR-1998; 98WO-CA00190.
 PR 25-FEB-1998; 98US-0030619.
 PR 10-MAR-1997; 97US-0040649.
 PR 20-AUG-1997; 97US-0915314.
 PR 26-SEP-1997; 97US-0060099.
 PA (MICR-) MICROLOGIX BIOTECH INC.
 PI Fraser JR, McNicol PJ, West MHP;
 DR WPI: 1998-520800/44.
 XX
 XX WPI: 1998-520800/44.
 PT New indolicidin peptide analogues - useful for, e.g. enhancing
 PT activity of antibiotic or overcoming tolerance, acquired resistance
 PT or inherent resistance of microorganisms
 PS
 SQ Claim 15; Page 93; 105pp; English.
 CC The present sequence represents a specifically claimed cationic peptide
 CC from the present invention. The present invention describes compositions
 CC and methods for treating infection, especially bacterial infections. The
 CC compositions and methods use cationic peptides in combination with an
 CC antibiotic agent which are then administered to a patient to enhance the
 CC activity of the antibiotic agent, to overcome: (a) tolerance; (b)
 CC acquired resistance; and (c) inherent resistance. The combinations of
 CC antibiotics and cationic peptides can provide synergistic activity
 CC against a microorganism that is tolerant, inherently resistant, or has
 CC acquired resistance to an antibiotic agent. They can be used for killing
 CC e.g. bacteria, fungi, parasites and viruses.
 XX
 SQ Sequence 21 AA:
 Query Match 100.0%; Score 86; DB 19; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KM leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KM multidrug resistance.
 OS Synthetic.
 PN WO9965506-A2.
 PD 23-DEC-1999.
 PF 14-JUN-1999; 99WO-CA00552.
 PR 12-JUN-1998; 98US-0096541.
 PA (MICR-) MICROLOGIX BIOTECH INC.
 PI Friedland HD, Krieger TJ, Taylor R, Effle D, Fraser JR, West MHP;
 DR WPI: 2000-223549/19.
 XX
 XX
 PT Novel pharmaceutical composition containing optionally activated
 PT polyoxalkylene-modified cationic peptides, useful for treating tumours
 PS
 SQ Disclosure: Page 15; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 21 AA:
 Query Match 100.0%; Score 86; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 10
 ID AAY91798 standard; Peptide: 21 AA.
 AC AAY91798;
 DT 06-JUN-2000 (first entry)
 DE Amino acid sequence of cationic peptide MBI 11B18CN.
 KM Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KM leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KM multidrug resistance.
 OS Synthetic.
 PN WO9965506-A2.
 PD 23-DEC-1999.
 PF 14-JUN-1999; 99WO-CA00552.
 PR 12-JUN-1998; 98US-0096541.

XX (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Friedland HD, Krieger TJ, Taylor R, Effle D, Fraser JR, West MHP;
 XX WPI; 2000-223549/19.
 DR
 XX
 PT Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS
 XX Disclosure; Page 15; 94pp; English.
 XX
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 21 AA;

Query Match 100.0%; Score 86; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 6.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPWMPWRRK 12
 |||||
 DB 1 ILRMPWMPWRRK 12

RESULT 11

AAM66363
 ID AAM66363 standard; peptide; 27 AA.

AC AAM66363;

DT 12-JAN-1999 (first entry)

DE Indolicidin analogue MBI 11B20.

KW Indolicidin analogue; resistance; cationic peptide; antibiotic;
 KW bacterial infection; tolerance; antibacterial; microorganism;
 KW bacteria; fungus; parasite; virus.

OS Bos taurus.

OS Synthetic.

PN WO9840401-A2.

PD 17-SEP-1998.

PF 10-MAR-1998; 98WO-CA00190.

PR 25-FEB-1998; 98US-0030619.

PR 10-MAR-1997; 97US-0040649.

PR 20-AUG-1997; 97US-0915314.

PR 26-SEP-1997; 97US-0060099.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Fraser JR, McNicol PJ, West MHP;

DR WPI; 1998-520800/44.
 PT New indolicidin peptide analogues - useful for, e.g. enhancing
 PT activity of antibiotic or overcoming tolerance, acquired resistance
 PT or inherent resistance of microorganisms
 XX

PS Claim 1; Page 91; 105pp; English.

XX
 CC The present sequence represents an indolicidin analogue. The present
 CC invention describes compositions and methods for treating infection,
 CC especially bacterial infections. The compositions and methods use
 CC cationic peptides in combination with an antibiotic agent which are
 CC then administered to a patient to enhance the activity of the antibiotic
 CC agent, to overcome: (a) tolerance; (b) acquired resistance; and (c)
 CC inherent resistance. The combinations of antibiotics and cationic
 CC peptides can provide synergistic activity against a microorganism that
 CC is tolerant, inherently resistant, or has acquired resistance to an
 CC antibiotic agent. They can be used for killing e.g. bacteria, fungi,
 CC parasites and viruses.
 XX

SQ Sequence 27 AA;

Query Match 100.0%; Score 86; DB 19; Length 27;
 Best Local Similarity 100.0%; Pred. No. 8.8e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPWMPWRRK 12
 |||||
 DB 1 ILRMPWMPWRRK 12

RESULT 12

AAV91800
 ID AAV91800 standard; Peptide; 28 AA.

AC AAV91800;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11B20CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.

OS Synthetic.

PN WO965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Effle D, Fraser JR, West MHP;

DR WPI; 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours

PS Claim 1; Page 15; 94pp; English.

XX
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX

SQ Sequence 28 AA:

Query Match 100.0%; Score 86; DB 21; Length 28;
 Best Local Similarity 100.0%; Pred. No. 9.1e-06;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPMPWRRK 12
 ||:|||||
 Db 1 ILRMPMPWRRK 12

RESULT 13

AAV24567

ID AAV24567 standard; peptide: 12 AA.

AC AAV24567;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #19.

KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
 KW additive; shampoo; soap; insecticide; herbicide; preservative;
 KW food; technical material.

OS Synthetic.

PN W09807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-US14779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;

DR WPI: 1998-169090/15.

PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid vectors, transformed cells and antibodies, also
 PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.

PS Claim 12; Page 89; 129pp; English.

XX AA24549 to AAV24615 represent indolicidin analogues of formulae
 CC (I)-(VIII) containing up to 25 amino acids (aa): RX2XX2B (I), BX2XX2B
 CC (II), BB2XX2XB (III), BX2XX2BBn(AA)nMLBBGS (IV), BX2XX2BB(AA)nM
 CC (V), LBBn2XX2XB (VI), LK2XX2XB (VII) and BB2XX2BB (VIII).
 CC Where 2 = P or V; X = hydrophobic residue, preferably W; B = basic aa,
 CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
 CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
 CC infections caused by bacteria (Gram positive or negative, or anaerobic);
 CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 CC trematodes) or viruses. Typical of very many pathogens that can be
 CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
 CC derived from the analogues may be used similarly; the compounds may
 CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
 CC may be used therapeutically or to coat medical devices; also they are
 CC useful as surface disinfectants, as additives to shampoo or soaps, as
 CC insecticides or herbicides, or as preservatives for foods and technical
 CC materials. The analogues are administered by injection, lavage, orally
 CC or topically, generally at 0.1-50 mg/Kg. These analogues have a broader
 CC spectrum of activity than indolicidin and modification as compounds
 CC reduces their toxicity.

XX SQ Sequence 12 AA:

Query Match 96.5%; Score 83; DB 19; Length 12;
 Best Local Similarity 91.7%; Pred. No. 1e-05;
 Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPMPWRRK 12
 ||:|||||
 Db 1 ILKMPMPWRRK 12

RESULT 14

AAV91788

ID AAV91788 standard; peptide: 12 AA.

AC AAV91788;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11B3CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.

OS Synthetic.

PN W0965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erle D, Fraser JR, West MHP;

DR WPI: 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PT .

PS Disclosure: Page 14; 94pp; English.

XX This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.

SQ Sequence 12 AA:

Query Match 96.5%; Score 83; DB 21; Length 12;
 Best Local Similarity 91.7%; Pred. No. 1e-05;
 Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRMPMPWRRK 12
 ||:|||||
 Db 1 ILKMPMPWRRK 12

RESULT 15

Search completed: January 4, 2002, 08:40:26
Job time: 109 sec

AAV24594

ID AAV24594 standard; peptide; 12 AA.

XX AAV24594;

XX 18-AUG-1999 (first entry)

XX Indolicidin analogue #46.

XX Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.

XX Synthetic.

XX W09807745-A2.

XX 26-FEB-1998.

XX 21-AUG-1997; 97WO-US14779.

XX 13-JAN-1997; 97US-0034949.

XX 21-AUG-1996; 96US-0024754.

XX (MICR-) MICROLOGIX BIOTECH INC.

XX Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;

XX WPI; 1998-169090/15.

XX New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.

XX Claim 14; Page 89; 129pp; English.

XX AAV24549 to AAV24615 represent indolicidin analogues of formulae
CC (I)-(VII) containing up to 25 amino acids (aa): RXZXXXB (I), BXZXXXB
CC (II), BBZXXZXXB (III), BXZXXZBBn(AA)nLBAGS (IV), BXZXXZBB(AA)nM
CC (V), LBnZXXnXXnXRK (VI), LKXZXXZXRK (VII) and BBZXXZBBB (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC derived from the analogues may be used similarly: the compounds may
CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
CC may be used therapeutically or to coat medical devices: also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.4-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.

XX Sequence 12 AA;

Query Match 95.3%; Score 82; DB 19; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.4e-05;

Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LRPWMPWRRK 12

DB 2 LRWMPWWRK 12

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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:57 ; Search time 24.75 Seconds
(without alignments)
10.911 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPRRK 12

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 212252 seqs, 22503292 residues

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : Issued Patents-AA:*

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4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/2/1aa/PCYUS.COMB.pep:*
6: /cgn2_6/ptodata/2/1aa/Backfile1.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	86	100.0	12	US-08-915-314-42	Sequence 42, Appl
2	86	100.0	20	US-08-915-314-47	Sequence 47, Appl
3	86	100.0	21	US-08-915-314-46	Sequence 46, Appl
4	86	100.0	21	US-08-915-314-48	Sequence 48, Appl
5	83	96.5	12	US-08-915-314-40	Sequence 40, Appl
6	82	95.3	12	US-08-915-314-76	Sequence 76, Appl
7	81	94.2	12	US-08-915-314-77	Sequence 77, Appl
8	81	94.2	12	US-08-915-314-78	Sequence 78, Appl
9	80	93.0	12	US-08-915-314-85	Sequence 85, Appl
10	80	93.0	12	US-08-915-314-86	Sequence 86, Appl
11	80	93.0	12	US-08-915-314-80	Sequence 80, Appl
12	78	90.7	12	US-08-915-314-83	Sequence 83, Appl
13	78	90.7	12	US-08-915-314-38	Sequence 38, Appl
14	78	90.7	12	US-08-915-314-69	Sequence 69, Appl
15	77	89.5	12	US-08-915-314-41	Sequence 41, Appl
16	75	87.2	12	US-08-915-314-52	Sequence 52, Appl
17	75	87.2	12	US-08-915-314-25	Sequence 25, Appl
18	75	87.2	13	US-08-915-314-30	Sequence 30, Appl
19	75	87.2	13	US-08-915-314-51	Sequence 51, Appl
20	75	87.2	13	US-08-915-314-62	Sequence 62, Appl
21	75	87.2	13	US-08-915-314-63	Sequence 63, Appl
22	75	87.2	13	US-08-915-314-64	Sequence 64, Appl
23	75	87.2	13	US-08-915-314-34	Sequence 34, Appl
24	75	87.2	13	US-08-702-054B-33	Sequence 33, Appl
25	75	87.2	13	US-08-702-054B-34	Sequence 34, Appl
26	75	87.2	13	US-08-702-054B-35	Sequence 35, Appl
27	75	87.2	13	US-09-042-071-36	Sequence 36, Appl

28	75	87.2	14	US-08-915-314-57	Sequence 57, Appl
29	75	87.2	16	US-08-702-054B-2	Sequence 2, Appl
30	75	87.2	21	US-08-915-314-54	Sequence 54, Appl
31	73	84.9	9	US-08-915-314-90	Sequence 90, Appl
32	73	84.9	11	US-08-915-314-44	Sequence 44, Appl
33	73	84.9	16	US-08-702-054B-38	Sequence 38, Appl
34	72	83.7	12	US-08-915-314-79	Sequence 79, Appl
35	72	83.7	12	US-08-915-314-81	Sequence 81, Appl
36	72	83.7	12	US-08-915-314-82	Sequence 82, Appl
37	72	83.7	12	US-08-915-314-84	Sequence 84, Appl
38	70	81.4	9	US-09-076-227-5	Sequence 5, Appl
39	70	81.4	10	US-09-076-227-4	Sequence 4, Appl
40	70	81.4	11	US-08-702-054B-9	Sequence 9, Appl
41	70	81.4	11	US-09-076-227-3	Sequence 3, Appl
42	70	81.4	12	US-08-915-314-39	Sequence 39, Appl
43	70	81.4	12	US-08-915-314-74	Sequence 74, Appl
44	70	81.4	12	US-08-702-054B-5	Sequence 5, Appl
45	70	81.4	12	US-09-076-227-2	Sequence 2, Appl

ALIGNMENTS

RESULT 1
US-08-915-314-42
; Sequence 42, Application US/08915314
; Patent No. 6180604
; GENERAL INFORMATION:
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Ertel, Douglas
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
; NUMBER OF SEQUENCES: 90
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/915,314
; FILING DATE: 20-AUG-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6180604/Leiburg Ph.D., Carol
; REGISTRATION NUMBER: 39,317
; REFERENCE/DOCKET NUMBER: 660081.405
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 42:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 12 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; US-08-915-314-42

Query Match 100.0%; Score 86; DB 4; Length 12;
Best Local Similarity 100.0%; Pred No. 1.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ILRPMWPRRK 12

DB 1 ILRWPMPWRRK 12

RESULT 2

US-08-915-314-47
Sequence 47, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 47:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-47

Query Match 100.0%; Score 86; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 2.2e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRRK 12
DB 1 ILRWPMPWRRK 12

RESULT 3
US-08-915-314-46
Sequence 46, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSEE: SEED and BERRY LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 46:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-46

Query Match 100.0%; Score 86; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILRWPMPWRRK 12
DB 1 ILRWPMPWRRK 12

RESULT 4
US-08-915-314-48
Sequence 48, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:

TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 48:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-48

Query Match 100.0%; Score 86; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 2.3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
DB 1 ILRPMWPMWRK 12

RESULT 5
US-08-915-314-40
Sequence 40, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-40

Query Match 96.3%; Score 83; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 3.5e-06;
Matches 11; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRPMWPMWRK 12
DB 1 ILRPMWPMWRK 12

RESULT 6
US-08-915-314-76
Sequence 76, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 76:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-76

Query Match 95.3%; Score 82; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 4.8e-06;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LRPMPWPMWRK 12
DB 2 LRPMPWPMWRK 12

RESULT 7
US-08-915-314-77
Sequence 77, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington

COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 77:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-77

Query Match 94.2%; Score 81; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 6.5e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRWPMPWRRK 12
| | | | | | | | | | | | | |
Db 1 IARWPMPWRRK 12

RESULT 8
US-08-915-314-87
Sequence 87, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Rife, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 87:

SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-87

Query Match 94.2%; Score 81; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 6.5e-06;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILRWPMPWRRK 11
| | | | | | | | | | | | | |
Db 1 ILRWPMPWRRK 11

RESULT 9
US-08-915-314-78
Sequence 78, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Rife, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 78:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-78

Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRWPMPWRRK 12
| | | | | | | | | | | | | |
Db 1 ILRWPMPWRRK 12

RESULT 10
US-08-915-314-85

Sequence 85, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 85:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-85

Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWMPWRRK 12
Db 1 ILRPMWMPWRRK 12

RESULT 11
US-08-915-314-86
Sequence 86, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 86:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-86

Query Match 93.0%; Score 80; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 9e-06;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWMPWRRK 12
Db 1 ILRPMWMPWRRK 12

RESULT 12
US-08-915-314-80
Sequence 80, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: NO. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 80:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid

STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-80

Query Match 90.7%; Score 78; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 1.7e-05;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWPMRRK 12
|||||
DB 1 ILRPMWPMRRK 12

RESULT 13
US-08-915-314-83
Sequence 83, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 83:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-83

Query Match 90.7%; Score 78; DB 4; Length 12;
Best Local Similarity 91.7%; Pred. No. 1.7e-05;
Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILRPMWPMRRK 12
|||||
DB 1 ILRPMWPMRRK 12

RESULT 14
US-08-915-314-38
Sequence 38, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:

APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 38:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-38

Query Match 90.7%; Score 78; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 1.9e-05;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 RMPWPMRRK 12
|||||
DB 4 RMPWPMRRK 13

RESULT 15
US-08-915-314-69
Sequence 69, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

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SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-69

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Query Match      89.58; Score 77; DB 4; Length 12;
Best Local Similarity 83.38; Pred. No. 2.3e-05;
Matches 10; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY      1 ILRPWMPWRRK 12
      1 :|||||||
Db      1 IKKPPWMPWRRK 12

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Search completed: January 4, 2002, 08:40:57
 Job time: 140 sec

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seq_documentation_block:
ID   AAA27298 standard; DNA; 114 BP.
XX
AC   AAA27298;
XX
DT   20-SEP-2000 (first entry)
XX
DE   Oligonucleotide used for synthesis of MBI 11B7 first cassette.
XX
KW   Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX   MBI-11; indolicidin; bovine; ss.
XX
OS   Synthetic.
XX
PN   WO200031279-A2.
XX
PD   02-JUN-2000.
XX
PE   19-NOV-1999; 99WO-CA01107.
XX
PR   20-NOV-1998; 98US-0109218.
XX
PA   (MICR-) MICROLOGIX BIOTECH INC.
XX
PI   Burian J, Bartfeld D;
XX
DR   WPI; 2000-400086/34.
XX
PT   Multi-domain fusion protein expression cassette used for high yield
PS   stable production of foreign peptide gene products -
XX
Example 5; Page 40; 73pp; English.
XX
CC   A novel method allows the efficient production of cationic peptides in
CC   recombinant host cells. The method involves construction of a
CC   multi-domain fusion protein expression cassette comprising a promoter and
CC   a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC   of anionic peptide sequences in the linker sequences neutralises the
CC   positive charge of the cationic peptide so that the charge of the
CC   fusion protein is controlled. This cassette allows high yield, stable
CC   production of the cationic peptide. Cationic peptides such as
CC   bovine indolicidin may be used as antimicrobial agents. The present
CC   sequence is an oligonucleotide that was used in the expression a
CC   cassette. MBI-11B7 is a cationic peptide derived from modifications
CC   of indolicidin.
XX
SQ   Sequence 114 BP; 20 A; 34 C; 32 G; 28 T; 0 other;

alignment_scores:
Quality: 86.00      Length: 12
Ratio: 7.167       Gaps: 0
Percent Similarity: 100.000  Percent Identity: 100.000

alignment_block:
US-09-444-281-36 x AAA27298

Align seg 1/1 to: AAA27298 from: 1 to: 114

1 11leuAgtTrrProTrrPrrParGArgLys 12
|||||
47 ATTCTGCGTTGGCGGTGGCGGTGCCGCAAA 82

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27294

seq_documentation_block:
ID   AAA27294 standard; DNA; 151 BP.
XX
AC   AAA27294;
XX
DT   20-SEP-2000 (first entry) ;
XX
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```
XX
DE   Oligonucleotide used for synthesis of MBI 2X11B7 last cassette.
XX
KW   Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX   MBI-11; indolicidin; bovine; ss.
XX
OS   Synthetic.
XX
PN   WO200031279-A2.
XX
PD   02-JUN-2000.
XX
PE   19-NOV-1999; 99WO-CA01107.
XX
PR   20-NOV-1998; 98US-0109218.
XX
PA   (MICR-) MICROLOGIX BIOTECH INC.
XX
PI   Burian J, Bartfeld D;
XX
DR   WPI; 2000-400086/34.
XX
PT   Multi-domain fusion protein expression cassette used for high yield
PS   stable production of foreign peptide gene products -
XX
Example 5; Page 38; 73pp; English.
XX
CC   A novel method allows the efficient production of cationic peptides in
CC   recombinant host cells. The method involves construction of a
CC   multi-domain fusion protein expression cassette comprising a promoter and
CC   a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC   of anionic peptide sequences in the linker sequences neutralises the
CC   positive charge of the cationic peptide so that the charge of the
CC   fusion protein is controlled. This cassette allows high yield, stable
CC   production of the cationic peptide. Cationic peptides such as
CC   bovine indolicidin may be used as antimicrobial agents. The present
CC   sequence is an oligonucleotide that was used to synthesise a
CC   cassette. MBI-11B7 is a cationic peptide derived from modifications
CC   of indolicidin.
XX
SQ   Sequence 151 BP; 22 A; 44 C; 49 G; 36 T; 0 other;

alignment_scores:
Quality: 86.00      Length: 12
Ratio: 7.167       Gaps: 0
Percent Similarity: 100.000  Percent Identity: 100.000

alignment_block:
US-09-444-281-36 x AAA27294

Align seg 1/1 to: AAA27294 from: 1 to: 151

1 11leuAgtTrrProTrrPrrParGArgLys 12
|||||
41 ATTCTGCGTTGGCGGTGGCGGTGCCGCAAA 76

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA1998.DAT:AAV60908

seq_documentation_block:
ID   AAV60908 standard; DNA; 88 BP.
XX
AC   AAV60908;
XX
DT   11-JAN-1999 (first entry)
XX
DE   DNA fragment encoding MB11.
XX
KW   MB128; cationic peptide; plasmid pRL1; small cryptic plasmid;
XX   replication; RepA; vector; RAMP; human; MB11; ss.
XX
OS   Synthetic.
```


OS Homo sapiens.
 XX
 PF MO9841636-A2.
 XX
 PD 24-SEP-1998.
 XX
 XX 16-MAR-1998; 98WO-CA00214.
 XX
 PR 14-MAR-1997; 97US-0040722.
 XX
 PA (BURT/) BURIAN J.
 PA (KAYW/) KAY W W.
 PI Burian J, Kay WW;
 XX
 DR WPI: 1998-531571/45.
 XX
 PT Increasing plasmid copy number in a cell with the repA gene product
 PT - and an small cryptic plasmid ori sequence, useful for high level
 PT expression of e.g. cytokines, antigens or therapeutic proteins
 XX
 PS Example 16; Page 57; 82pp; English.
 XX
 CC This oligonucleotide was used as a template in a PCR reaction (see
 CC also AAV60909-10) to generate a DNA fragment encoding the cationic
 CC peptide MB11 (see AAV71690). The PCR product was cloned into
 CC the universal vector pR2h-B1, which contains the R21 replication
 CC leader of RepA (see AAV71686) and 2 tandem copies of the prepro
 CC region (hpro) of human defensin. The vector provides expression
 CC of R21-hpro-MB11 fusion in host cells. The invention provides a
 CC controlled replication plasmid vectors (RAMP vectors) comprising a
 CC replication origin of a small cryptic plasmid such as pK11 (see
 CC AAV58292) and a gene encoding RepA (see AAV71686). The vectors can
 CC reach very high levels of plasmid replication, but are not lethal
 CC to the host cell, and can be used to direct the high level
 CC expression of e.g. cytokines, antigens and therapeutic proteins.
 XX
 SQ Sequence 88 BP; 20 A; 18 C; 25 G; 25 T; 0 other;

alignment_scores:
 Quality: 75.00 Length: 10
 Ratio: 7.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:
 US-09-444-281-36 x AAV60908 ..

Align seg 1/1 to: AAV60908 from: 1 to: 88

3 ArgTrpProTrpTrpProTrpArgLys 12
 :::::::::::::::::::::::::::::::
 34 AATGCGCGTGTGTGCGCGCGTGTGTANA 63

seq_name: /SIDS8/gcgdata/geneseq/geneseq/NA2000.DAT:AAA27291

seq_documentation_block:
 ID AAA27291 standard; DNA; 114 BP.
 XX
 AC AAA27291;
 XX
 DT 20-SEP-2000 (first entry)
 XX
 DE Oligonucleotide used for synthesis of MB1-11 fragment.
 XX
 KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
 KW MB1-11; indolicidin; bovine; ss.
 XX
 OS Synthetic.
 OS
 PN WO200031279-A2.
 PD 02-JUN-2000.

XX
 XX 19-NOV-1999; 99WO-CA01107.
 XX
 PR 20-NOV-1998; 98US-0109218.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 PA
 PI Burian J, Bartfeld D;
 XX
 DR WPI: 2000-400086/34.
 XX
 PT Multi-domain fusion protein expression cassette used for high yield
 PT stable production of foreign peptide gene products -
 XX
 PS Example 4; Page 37; 73pp; English.
 XX
 CC A novel method allows the efficient production of cationic peptides in
 CC recombinant host cells. The method involves construction of a
 CC multi-domain fusion protein expression cassette comprising a promoter and
 CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
 CC of anionic peptide sequences in the linker sequences neutralises the
 CC positive charge of the cationic peptide so that the charge of the
 CC fusion protein is controlled. This cassette allows high yield, stable
 CC production of the cationic peptide. Cationic peptides such as
 CC bovine indolicidin may be used as antimicrobial agents. The present
 CC sequence is an oligonucleotide that was used to synthesise a
 CC MB1-11 fragment. MB1-11 is a cationic peptide derived from modifications
 CC of indolicidin.
 XX
 SQ Sequence 114 BP; 25 A; 26 C; 30 G; 33 T; 0 other;

alignment_scores:
 Quality: 75.00 Length: 10
 Ratio: 7.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:
 US-09-444-281-36 x AAA27291 ..

Align seg 1/1 to: AAA27291 from: 1 to: 114

3 ArgTrpProTrpTrpProTrpArgLys 12
 :::::::::::::::::::::::::::::::
 50 AATGCGCGTGTGTGCGCGCGTGTGTANA 79

seq_name: /SIDS8/gcgdata/geneseq/geneseq/NA1999.DAT:AAV83788

seq_documentation_block:
 ID AAV83788 standard; DNA; 39 BP.
 XX
 AC AAV83788;
 XX
 DT 19-MAR-1999 (first entry)
 XX
 DE Antimicrobial peptide Indolicidin encoding DNA.
 XX
 KW Antimicrobial; fusion; acidic peptide; recombinant; microorganism;
 KW guamerin; basic peptide; indolicidin; ss.
 XX
 OS Synthetic.
 OS Bos sp.
 FH Key Location/Qualifiers
 FT CDS 1..39 /*tag= a
 FT /note= "the start and stop codons are not indicated"
 XX
 PN WO9854336-A1.
 PD 03-DEC-1998.
 XX
 PF 28-MAY-1998; 98WO-KR00132.

```
XX 09-APR-1998; 98KR-0013372.
PR 28-MAY-1997; 97KR-0021312.
XX
XX (KOAD ) KOREA ADV INST SCI & TECHNOLOGY.
PA (SAMY-) SAMYANG GENEX CORP.
XX
XX Hong S, Kang MH, Kim JH, Kim S, Lee H, Lee JH;
PI WPI; 1999-059844/05.
DR P-PSDB; AAW87609.
XX
XX New method for mass production of antimicrobial peptides - by
PT constructing fusion genes comprising acidic and antimicrobial
PT peptide genes and transforming host with vector containing these
XX
XX Example 6; Page 18; 52pp; English.
XX
XX The invention relates to mass production of antimicrobial peptides. The
CC method comprises constructing a fusion gene containing a first gene
CC encoding a negatively charged acidic peptide having at least two cysteine
CC residues, and a second gene encoding a positively charged basic
CC antimicrobial peptide. A host microorganism is transformed with a vector
CC containing the fusion gene and then cultured. The expressed antimicrobial
CC peptide is then recovered. The method is used to mass produce
CC antimicrobial peptides in recombinant microorganisms. The inhibitory
CC effect of the expressed antimicrobial peptide upon the growth of the host
CC microorganism is considerably reduced by fusing it to the acidic peptide.
CC Therefore, the use of the fusion gene provides an economic, recombinant
CC alternative of mass producing antimicrobial peptides, which overcomes the
CC disadvantages of low-productivity and poor economy, previously
CC encountered by recombinant and chemical methods. The present sequence
CC represents the DNA encoding an antimicrobial peptide indolicidin. This
CC can be used along with the acidic peptide Guamerin gene in the
CC construction of the fusion gene.
XX
SQ Sequence 39 BP; 4 A; 10 C; 16 G; 9 T; 0 other;

alignment_scores:
      Quality: 70.00      Length: 9
      Ratio: 7.778      Gaps: 0
Percent Similarity: 100.000      Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAV83788 ..

Align seq 1/1 to: AAV83788 from: 1 to: 39
      3 ArgTrpProTrpTrpArgArg 11
      ::::::::::::::::::::11
      13 AATGCGCGTGTGCGCGTGTCT 39

seq_name: /SID58/gcgdata/geneseq/NA2000.DAT:AAZ29389

seq_documentation_block:
ID AAZ29389 standard; DNA; 47 BP.
XX
XX AAZ29389;
XX
XX 29-FEB-2000 (first entry)
XX
XX PCR primer-15 for synthesis of antimicrobial peptide indolicidin gene.
DE
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
XX Synthetic.
XX
XX WO964611-A1.
XX
```

```
PD 16-DEC-1999.
XX
XX 08-JUN-1999; 99WO-KR00282.
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
XX
XX Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
PI WPI; 2000-097542/08.
DR
XX
XX New DNA constructs useful for mass production of antimicrobial peptides
PT in microorganism hosts -
PT
XX
XX Example 1; Page 13; 67pp; English.
XX
XX The present sequence is a chemically synthesised PCR primer which was
CC used to synthesise a gene encoding antimicrobial peptide indolicidin.
CC The antimicrobial peptide gene is used in a DNA construct that comprises
CC entire, partial or a derivative of purf gene (glutamine
CC pyrophosphoribosyl pyrophosphate amidotransferase gene). The DNA
CC construct allows mass production of the antimicrobial peptide in
CC microbial hosts without killing the host cells. The antimicrobial
CC peptides are useful commercially in the pharmaceutical and
XX food industries.
XX
SQ Sequence 47 BP; 6 A; 11 C; 19 G; 11 T; 0 other;

alignment_scores:
      Quality: 70.00      Length: 9
      Ratio: 7.778      Gaps: 0
Percent Similarity: 100.000      Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AAZ29389 ..

Align seq 1/1 to: AAZ29389 from: 1 to: 47
      3 ArgTrpProTrpTrpArgArg 11
      ::::::::::::::::::::11
      17 AATGCGCGTGTGCGCGTGTCT 43

seq_name: /SID58/gcgdata/geneseq/NA2000.DAT:AAZ29390

seq_documentation_block:
ID AAZ29390 standard; DNA; 47 BP.
XX
XX AAZ29390;
XX
XX 29-FEB-2000 (first entry)
XX
XX PCR primer-16 for synthesis of antimicrobial peptide indolicidin gene.
DE
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
XX Synthetic.
XX
XX WO964611-A1.
XX
XX 16-DEC-1999.
XX
XX 08-JUN-1999; 99WO-KR00282.
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
XX
```

```

PI  Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX
XX  Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
DR  WPI: 2000-097542/08.
XX
XX  New DNA constructs useful for mass production of antimicrobial peptides
PT  in microorganism hosts
XX
XX  Example 1; Page 13; 67pp; English.
PS
CC  The present sequence is chemically synthesised PCR primer which was
CC  used to synthesise a gene encoding antimicrobial peptide Indolicidin.
CC  The antimicrobial peptide gene is used in a DNA construct that comprises
CC  entire, partial or a derivative of purf gene (glutamine
CC  pyrophosphoribosyl pyrophosphate amidotransferase gene). The DNA
CC  construct allows mass production of the antimicrobial peptide in
CC  microbial hosts without killing the host cells. The antimicrobial
CC  peptides are useful commercially in the pharmaceutical and
CC  food industries.
XX
S0  Sequence 47 BP; 12 A; 18 C; 10 G; 7 T; 0 other;
XX
XX
XX  Alignment scores:
XX      Quality: 70.00      Length: 9
XX      Ratio: 7.778      Gaps: 0
XX  Percent Similarity: 100.000      Percent Identity: 88.889
XX
XX  alignment_block:
XX  US-09-444-281-36 x AA229390/rev ..
XX
XX  Align seg 1/1 to reverse of: AA229390 from: 1 to: 47
XX
XX      3 ATGTTPPOTPTPTPPTCTPARGTG 11
XX      ::::::::::::::::::::::::::::
XX      35 AATGCGCCGTGCTGCGCCTGCGCTGT 9
XX
XX  seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2000.DAT:AA229364
XX
XX  seq_documentation_block:
XX  ID AA229364 standard; DNA; 53 BP.
XX
XX  AC AA229364;
XX
XX  DT 29-FEB-2000 (first entry)
XX
XX  DE Antimicrobial peptide, Indolicidin encoding DNA.
XX
XX  purf gene: glutamine pyrophosphoribosyl pyrophosphate amidotransferase;
XX  purf derivative; fusion partner; antimicrobial peptide; indolicidin;
XX  mass production; cleavage site; hydroxylamine; CNBr; DNA construct; cow;
XX  neutralise; toxicity; pharmaceutical industry; food industry; ds.
XX
XX  Bos taurus.
XX
XX  OS
XX  FH Key Location/Qualifiers
XX  FT CDS 5..46
XX  FT /*tag= a
XX  FT /*product= "Indolicidin peptide"
XX  FT /note= "Antimicrobial peptide used in DNA construct"
XX
XX  WO9964611-A1.
XX
XX  PD 16-DEC-1999.
XX
XX  PE 08-JUN-1999; 99WO-KR00282.
XX
XX  PR 09-JUN-1998; 98KR-0022117.
XX  PR 14-MAY-1999; 99KR-0017920;
XX
XX  RA (SAMT-) SAMYANG GENEX CORP.
XX
XX  Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;

```

```

XX  WPI: 2000-097542/08.
DR  P-PSDB: AA144324.
XX
XX  New DNA constructs useful for mass production of antimicrobial peptides
PT  in microorganism hosts -
XX
XX  Claim 1; Fig 1; 67pp; English.
XX
XX  The present DNA sequence encodes an antimicrobial peptide, Indolicidin
CC  derived from cow, Bos taurus. It is used along with a derivative
CC  of pufE gene sequence that functions as a fusion partner.
CC  A DNA construct that comprises, this antimicrobial peptide encoding
CC  sequence and the entire, partial or derivative of pufE gene, is used for
CC  mass production of the antimicrobial peptide in microorganisms without
CC  killing the host cells. Use of the pufE gene derivative sequence,
CC  neutralises the toxicity of the antimicrobial peptides against the host
CC  microorganism. The antimicrobial peptides are useful commercially in the
CC  pharmaceutical and food industries.
XX
XX  Sequence 53 BP; 8 A; 12 C; 20 G; 13 T; 0 other;
XX
XX  alignment_scores:
XX      Quality: 70.00      Length: 9
XX      Ratio: 7.778      Gaps: 0
XX      Percent Similarity: 100.000      Percent Identity: 88.889
XX
XX  alignment_block:
XX  US-09-444-281-36 x AA229364
XX
XX  Align seg 1/1 to: AA229364 from: 1 to: 53
XX
XX      3 ArgTrpProTyrPTrProTyrPArg 11
XX      ::::::::::::::::::::::::::::
XX      17 AAATGGCCGTGCTGCCCTGGCGTCTGT 43
XX
XX  seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2000.DAT:AA240246
XX
XX  seq_documentation_block:
XX  ID AA240246 standard; DNA: 69 BP.
XX
XX  AC AA240246;
XX
XX  DT 23-FEB-2000 (first entry)
XX
XX  DE Oligonucleotide for cloning indolicidin peptide coding sequence.
XX
XX  KW Indolicidin; bacitracin; sulphate-reducing bacteria; growth inhibitor;
XX  corrosion; degradation; metal; concrete; cement; dental implant; biofilm;
XX  ss.
XX
XX  OS Synthetic.
XX
XX  OS Bacillus sp.
XX
XX  PN W09956553-A1.
XX
XX  PD 11-NOV-1999.
XX
XX  PE 03-MAY-1999; 99MO-US09675.
XX
XX  PR 06-MAY-1998; 98US-0074037.
XX  PR 31-MAR-1999; 99US-0282277.
XX
XX  PA (REGC ) UNIV CALIFORNIA.
XX
XX  PI Wood TK, Jayaraman A, Earhman JC;
XX
XX  DR WPI: 2000-052882/04.
XX
XX  PT Inhibiting growth of sulphate-reducing bacteria using other bacteria,
XX  particularly for protection of metals and concrete -
XX

```

PS Example 4; Fig 1; 84pp; English.
 CC This sequence represents an oligonucleotide for cloning the non-ramified
 CC indolicidin peptide coding sequence. The invention relates to a method
 CC for inhibiting growth of sulphate-reducing bacteria (A) on a material (B)
 CC sensitive to corrosion or degradation, by applying to (B) a bacterium (C)
 CC that secretes a compound (I) able to inhibit growth of (A). The method is
 CC used to protect metal, concrete or cement against corrosion and
 CC degradation, but (B) can also be used to protect dental implants. (B) is
 CC present in an open or closed system (e.g. water cooling tower, liquid
 CC storage container, fuel tank, sewer or drainage system etc.) or part of a
 CC bridge or other structure. The method is more effective and less
 CC expensive than known methods for inhibiting (A), and reduces the amount
 CC of toxic chemicals released. Conventional biofilms of aerobic organisms
 CC tend to encourage growth of (A), and addition of (C) to the biofilm
 CC prevents this. A single application of (C) lasts for a long time, and (I)
 CC are produced exactly where they are required and inhibit (A) without
 CC significant impact on other organisms (this effect includes reducing
 CC resistance of (A) to conventional biocides, which may then be used in
 CC reduced amounts). If local damage to the biofilm occurs, the underlying
 CC material is still protected by diffusion of (I) from neighbouring areas.
 CC
 SQ Sequence 69 BP; 14 A; 18 C; 20 G; 17 T; 0 other;

alignment_scores:
 Quality: 70.00 Length: 9
 Ratio: 7.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
 US-09-444-281-36 x AAZ40246 ..

Align seg 1/1 to: AAZ40246 from: 1 to: 69

```

3 ArgTrpProTrpTrpProTrpArgArg 11
:::|||||
28 AAATGGCCTGTGTGGCTTGGCGCGC 54

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seq_name: /SIDSB8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ49764

seq_documentation_block:
 ID AAZ49764 standard; DNA; 211 BP.

AC AAZ49764;

DT 18-APR-2000 (first entry)

DE Poly-(Indol (1-13)-Met-Ala-Arg-Ile-Ala-Met)3 DNA.

XX Crosslinked indolicidin analog; X-Indolicidin; poly-Indol 1-13;
 KW stability; bovine neutrophil; antimicrobial; antibacterial; fungicide;
 KW protozoacide; virucide; anti-HIV; human immunodeficiency virus-1;
 KW HIV-1; gram positive bacteria; gram negative; Staphylococcus aureus;
 KW Escherichia coli; Salmonella typhimurium; yeast; fungi; protozoa;
 KW Candida albicans; Cryptococcus neoformans; Giardia; Acanthamoeba;
 KW hexapeptide spacer; ds.

XX Synthetic.

OS Bos sp.

Key Location/Qualifiers
 CDS 8..199

FT /tag= a
 FT /product= "Poly-(Indol(1-13)-Met-Ala-Arg-Ile-Ala-Met)3"
 FT /note= "encodes three copies of Indol 1-13, each
 FT separated by Met-Ala-Arg-Ile-Ala-Met spacer sequence"

FT primer_bind
 FT 1..21

FT complement (191..211)
 FT /tag= b

FT primer_bind
 FT /tag= c

FT misc_feature
 FT 68..71
 FT /tag= d

FT /note= "corresponds to overlap in oligonucleotides
 FT used for ligation"
 FT 148..151
 FT misc_feature
 FT /tag= e
 FT /note= "corresponds to overlap in oligonucleotides
 FT used for ligation"
 PN W09965510-A1.
 PD 23-DEC-1999.
 XX
 XX 20-MAY-1999; 99MO-US11165.
 XX
 XX 18-JUN-1998; 98US-0099631.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 XX Selsled ME, Osapay K;
 PI
 XX
 DR WPI: 2000-147133/13.
 DR P-PSDB; AAY44668.

XX Crosslinked indolicidin analogs with antimicrobial activity against
 PT bacteria, yeast, fungi, protozoa and viruses
 XX
 PS Example 1C; Fig 1; 53pp; English.

CC The patent discloses crosslinked analogs of indolicidin (indol 1-13)
 CC which is a naturally occurring peptide isolated from bovine neutrophils
 CC and has antimicrobial activity. The crosslinked indolicidin
 CC (X-Indolicidin) analogs are stable and have antimicrobial activity
 CC against gram positive and negative bacteria (e.g. Staphylococcus aureus,
 CC Escherichia coli and Salmonella typhimurium), yeasts and fungi (e.g.
 CC Candida albicans, Cryptococcus neoformans), protozoa (e.g. Giardia
 CC species and Acanthamoeba species), and viruses (e.g. HIV-1).
 CC They can be used for reducing or inhibiting the growth or survival of
 CC microorganisms in an environment e.g. a food or food product, a
 CC solution, an inanimate object comprising a surface, or a mammal.
 CC The present sequence is a DNA encoding a protein comprising three
 CC copies of indol 1-13 each separated by a hexapeptide spacer sequence.
 CC The sequence was used to produce a recombinant construct for the
 CC expression of indol-homoserine (Hse) analog. The ability of
 CC indol-Hse analog to maintain antimicrobial activity provides a means to
 CC produce X-Indolicidin analog precursors in sufficient quantities.

XX Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
 Quality: 70.00 Length: 9
 Ratio: 7.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:
 US-09-444-281-36 x AAZ49764 ..

Align seg 1/1 to: AAZ49764 from: 1 to: 211

```

3 ArgTrpProTrpTrpProTrpArgArg 11
:::|||||
38 AAATGGCCTGTGTGGCTTGGCGCTGT 64

```

seq_name: /SIDSB8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ45123

seq_documentation_block:
 ID AAZ45123 standard; DNA; 211 BP.

AC AAZ45123;

DT 28-FEB-2000 (first entry)

XX Indolicidin fusion peptide nucleotide sequence.

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:26 : Search time 53.46 Seconds
(without alignments)
18.013 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKMPMPWRRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues
Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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22: /SIDSB/gcgdata/geneseq/geneseq/AA2000.DAT.*
23: /SIDSB/gcgdata/geneseq/geneseq/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	91	100.0	13	18 AAW12873	Antimicrobial catl
2	91	100.0	13	19 AAY24569	Indolicidin analog
3	91	100.0	13	19 AAW6378	Cationic peptide o
4	91	100.0	13	19 AAW71690	Cationic peptide M
5	91	100.0	13	21 AAY94495	MBI-11 peptide der
6	91	100.0	13	21 AAY92795	Indolicidin analog
7	91	100.0	13	21 AAY91773	Amino acid sequenc
8	91	100.0	13	21 AAY91774	Amino acid sequenc
9	91	100.0	13	21 AAY91818	Amino acid sequenc
10	91	100.0	13	21 AAY91819	Amino acid sequenc
11	91	100.0	13	21 AAY91820	Amino acid sequenc

12	91	100.0	14	19 AAY24583	Indolicidin analog
13	91	100.0	14	21 AAY91811	Amino acid sequenc
14	91	100.0	21	19 AAY24582	Indolicidin analog
15	91	100.0	21	21 AAY91806	Amino acid sequenc
16	87	95.6	12	19 AAY24580	Indolicidin analog
17	87	95.6	12	21 AAY91804	Amino acid sequenc
18	86	94.5	12	18 AAW12877	Antimicrobial catl
19	86	94.5	12	19 AAY24615	Indolicidin analog
20	86	94.5	12	21 AAY91833	Amino acid sequenc
21	86	94.5	13	19 AAY24572	Indolicidin analog
22	86	94.5	13	21 AAY91812	Amino acid sequenc
23	86	94.5	14	21 AAY24573	Indolicidin analog
24	86	94.5	14	21 AAY91813	Amino acid sequenc
25	86	94.5	15	18 AAW13802	Antimicrobial catl
26	86	94.5	20	19 AAY24570	Indolicidin analog
27	86	94.5	20	21 AAY91807	Amino acid sequenc
28	86	94.5	21	19 AAY24571	Indolicidin analog
29	86	94.5	21	21 AAY91808	Amino acid sequenc
30	85	93.4	12	19 AAY24586	Indolicidin analog
31	85	93.4	12	21 AAY91828	Amino acid sequenc
32	85	93.4	13	18 AAW27179	Antimicrobial catl
33	85	93.4	13	18 AAW12869	Antimicrobial catl
34	85	93.4	13	18 AAW12894	Antimicrobial catl
35	85	93.4	13	19 AAY24610	Indolicidin analog
36	85	93.4	13	19 AAY24565	Indolicidin analog
37	85	93.4	13	21 AAY91786	Amino acid sequenc
38	85	93.4	13	21 AAY91795	Amino acid sequenc
39	83	91.2	12	19 AAY24568	Indolicidin analog
40	83	91.2	12	21 AAY91789	Amino acid sequenc
41	83	91.2	13	18 AAW12892	Antimicrobial catl
42	83	91.2	13	18 AAW12893	Antimicrobial catl
43	83	91.2	13	18 AAW12896	Antimicrobial catl
44	83	91.2	13	18 AAW12897	Antimicrobial catl
45	83	91.2	13	19 AAY24612	Indolicidin analog

ALIGNMENTS

RESULT 1
AAW12873
ID AAW12873 standard; peptide; 13 AA.
AC AAW12873:
XX
XX 10-DEC-1997 (first entry)
DT
XX
XX Antimicrobial cationic peptide CP-11.
DE
XX
XX Bacterial; viral; antitumour; food; preservative; inhibitor; growth;
KW Bacterium; yeast; endotoxaemia; sepsis; antibiotic; fungal;
KW antiviral; candida albicans; steriliant; Salmonella; Yersinia;
KW Shigella.
OS
XX Synthetic.
XX
XX W09708199-A2.
XX
XX 06-MAR-1997.
XX
XX 23-AUG-1996; 96WO-IB00996.
XX
XX 23-AUG-1995; 95US-0002687.
XX
XX (UYBR-) UNIV BRITISH COLUMBIA.
XX
XX Falla TJ, Cough M, Hancock RW;
XX
XX WPI: 1997-179179/16.
XX
XX Cationic peptide(s) having anti-microbial activity - used for the
XX inhibition of bacterial and viral growth, as an antitumour agent,
XX and as a food preservative
PT

XX Claim 2; Page 65; 89pp; English.

XX The present sequence represents a specifically claimed novel isolated
XX cationic peptide which has antimicrobial activity. The amino acid
XX sequence of antimicrobial cationic peptides (including the present
XX sequence) is selected from: X1X1ProX2X3X2Pro(X2X2Pro)X2X3(X5)O;
XX X1X1ProX2X3X4(X5)ProX2X3X3; X1X1X3(ProTrp)uX3X2X5X2X5X2(X5)O;
XX X1X1X3X3X2Pro(X2X2Pro)X2(X5)m; where m = 1-5; n = 1-2; o = 2-5; r
XX = 0-8; u = 0-1; X1 = Ile, Leu, Val, Phe, Tyr, Trp or Met; X2 = Trp or
XX Phe; X3 = Arg or Lys; X4 = Trp or Lys; and X5 = Phe, Trp, Arg, Lys or
XX Pro. The peptides are preferably amidated or carboxymethylated. The
XX peptides may be used in methods for inhibiting the growth of a bacterium
XX or yeast, or for inhibiting an endotoxaemia or sepsis associated
XX disorder in a subject. The peptides have a broad activity against
XX antibiotic resistant bacteria, combined with activity against the
XX medically important fungus *Candida albicans*. In addition, the peptides
XX are useful as antileukemia agents and/or antiviral agents. The peptides
XX may be used as sterilants or preservatives of materials susceptible to
XX microbial or viral contamination, e.g. in processed foods to inhibit
XX *Salmonella*, *Yersinia* and *Shigella*. The peptides are compact and tend to
XX have a unique polypyrrolone type II extended helix structure that permits
XX them to span the membrane with relatively few amino acids. The peptides
XX possess the ability to work synergistically with antibiotics, and in
XX addition, some of them possess anti-endotoxin activity.
XX N.B. The present sequence represents SEQ ID NO:1 in the claims and
XX examples of the specification, but differs slightly from the SEQ ID NO:1
XX in the sequence listing on page 51 of the specification (see AAW271179).

PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxyalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.
XX
XX Example 1; Page 32; 129pp; English.
PS
XX AAY24549 to AAY24615 represent indolicidin analogues of formulae
XX (I)-(VIII) containing up to 25 amino acids (aa): RZXXZXB (I), BZXZXZB
XX (II), BBRXZXZXZB (III), BXZXZXBBn(AA)nMILBACS (IV), BXZXZXBB(AA)nM
XX (V), LBBnXZXZXZB (VI), LKXZXZXZB (VII) and BBRXZXZXBBB (VIII).
XX Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
XX preferably R or K; AA = any aa, n = 0 or 1; in (II), at least 1 Z = V;
XX in (VIII) at least 2 X = F or Y. The analogues are used to treat
XX infections caused by bacteria (Gram positive or negative, or anaerobic);
XX fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
XX trematodes) or viruses. Typical of very many pathogens that can be
XX controlled are *Leishmania*, *Trypanosoma*, *Ascaris lumbricoides*, *Fasciola*
XX *hepatica*, *Klebsiella pneumoniae*, *Bordetella pertussis*, *Staphylococcus*
XX *aureus*, *Listeria*, *Clostridium*, *rotavirus* and *papilloma virus*. Compounds
XX derived from the analogues may be used similarly; the compounds may
XX also be prepared from antibiotics or antiparasitic agents. The analogues
XX may be used therapeutically or to coat medical devices; also they are
XX useful as surface disinfectants, as additives to shampoo or soaps, as
XX insecticides or herbicides, or as preservatives for foods and technical
XX materials. The analogues are administered by injection, lavage, orally
XX or topically, generally at 0.1-50 mg/kg. These analogues have a broader
XX spectrum of activity than indolicidin and modification as compounds
XX reduces their toxicity.

SO Sequence 13 AA;

Query Match 100.0%; Score 91; DB 18; Length 13;
Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
Db 1 ILKKWPMWPMRRK 13

RESULT 2

AAY24609
ID AAY24609 standard; peptide; 13 AA.

AC AAY24609;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #61.

XX Indolicidin: bacterial infection; photo-oxidised solubilisier;
XX antimicrobial; antibiotic; antiarrhythmic; surface disinfectant;
XX additive; shampoo; soap; insecticide; herbicide; preservative;
XX food; technical material.

OS Synthetic.

PN WO9807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-US14779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Erle D, Fraser JR, Krieger TJ, Taylor R, West MH;

DR WPI; 1998-169090/15.

Query Match 100.0%; Score 91; DB 19; Length 13;
Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
Db 1 ILKKWPMWPMRRK 13

RESULT 3

AAW6378
ID AAW6378 standard; peptide; 13 AA.

AC AAW6378;

DT 12-JAN-1999 (first entry)

DE Cationic peptide of claim 15 #5.

XX Indolicidin analogue; resistance; cationic peptide; antibiotic;
XX bacterial infection; tolerance; antibacterial; microorganism;
XX bacteria; fungus; parasite; virus.

OS Synthetic.

PN WO9840401-A2.

PD 17-SEP-1998.

PF 10-MAR-1998; 98WO-CA00190.

PR 25-FEB-1998; 98US-0030619.

PR 10-MAR-1997; 97US-0040649.

PR 20-AUG-1997; 97US-0915314.

PR 26-SEP-1997; 97US-0060099.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Fraser JR, McNicol PJ, West MHP;

DR WPI: 1998-520800/44.
XX New indolicidin peptide analogues, useful for, e.g. enhancing
PT activity of antibiotic or overcoming tolerance, acquired resistance
PT or inherent resistance of microorganisms
XX
PS Claim 15; Page 93; 105pp; English.
XX
CC The present sequence represents a specifically claimed cationic peptide
CC from the present invention. The present invention describes compositions
CC and methods for treating infection, especially bacterial infections. The
CC compositions and methods use cationic peptides in combination with an
CC antibiotic agent which are then administered to a patient to enhance the
CC activity of the antibiotic agent, to overcome: (a) tolerance; (b)
CC acquired resistance; and (c) inherent resistance. The combinations of
CC antibiotics and cationic peptides can provide synergistic activity
CC against a microorganism that is tolerant, inherently resistant, or has
CC acquired resistance to an antibiotic agent. They can be used for killing
CC e.g. bacteria, fungi, parasites and viruses.
XX
SQ Sequence 13 AA;

Query Match 100.0%; Score 91; DB 19; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
DB 1 ILKKWPMWPMRRK 13

RESULT 4
AAW71690
ID AAW71690 standard; Peptide: 13 AA.
XX
AC AAW71690;
XX
DT 11-JAN-1999 (first entry)
XX
DE Cationic peptide MB111 (MW 1879).
XX
KM MB111; cationic peptide; plasmid pK11; small cryptic plasmid;
KW replication; RepA; vector; RAMP.
XX
OS Synthetic.
XX
PN WO9841636-A2.
XX
PD 24-SEP-1998.
XX
PE 16-MAR-1998; 98MO-CA00214.
XX
PR 14-MAR-1997; 97US-0040722.
XX
PA (BURI/) BURIAN J.
PA (KAYW/) KAY W W.
XX
PI Burian J, Kay WW;
XX
DR WPI: 1998-531571/45.
XX
PT Increasing plasmid copy number in a cell with the repA gene product
PT - and an small cryptic plasmid ori sequence, useful for high level
PT expression of e.g. cytokines, antigens or therapeutic proteins
XX
XX Example 13; Page 54; 82pp; English.
XX
CC MB111 is a small (mol. wt. 1879) cationic peptide. DNA encoding
CC MB111 has been incorporated into vector pK11-B1, in which the
CC replication leader (R21) sequence of RepA (see also AAW71686) is
CC joined to 2 Hpro peptides (see also AAW71692), to provide a
CC vector for expression of MB111 in host cells. The invention

CC provides controlled replication plasmid vectors (RAMP vectors)
CC comprising a replication origin of a small cryptic plasmid and a
CC gene encoding RepA. The vectors can reach very high levels of
CC plasmid replication, but are not lethal to the host cell, and can
CC be used to direct the high level expression of e.g. cytokines,
CC antigens and therapeutic proteins.
XX
SQ Sequence 13 AA;

Query Match 100.0%; Score 91; DB 19; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13
DB 1 ILKKWPMWPMRRK 13

RESULT 5
AA94495
ID AA94495 standard; Peptide: 13 AA.
XX
AC AA94495;
XX
DT 20-SEP-2000 (first entry)
XX
DE MB1-11 peptide derived from indolicidin.
XX
KM Cellulose binding domain; CBD; cationic peptide;
KM MB1-11; indolicidin; bovine.
XX
OS Bos taurus.
XX
PN WO200031279-A2.
XX
PD 02-JUN-2000.
XX
PE 19-NOV-1999; 99MO-CA01107.
XX
PR 20-NOV-1998; 98US-0109218.
XX
PA (MICR-) MICROLOGIX BIOTECH INC.
XX
PI Burian J, Bartfeld D;
XX
DR WPI: 2000-400086/34.
XX
PT Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
XX
PS Disclosure; Page 24; 73pp; English.
XX
CC A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is the MB1-11 peptide. MB1-11 is a cationic peptide derived
CC from modifications of indolicidin.
XX
SQ Sequence 13 AA;

Query Match 100.0%; Score 91; DB 21; Length 13;
Best Local Similarity 100.0%; Pred. NO: 7.2e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKWPMWPMRRK 13

Db 1 ILKWPMPWRK 13

RESULT 6

ID AAY92795 standard; peptide; 13 AA.

AC AAY92795;

DT 29-AUG-2000 (first entry)

DE Indolicidin analogue, CP-11.

KM Magainin; antimicrobial; transgenic plant; protease degradation; Rev4;

KW Indolicidin; protein production; reverse peptide.

OS Synthetic.

PN WO200026344-A1.

PD 11-MAY-2000.

PF 29-OCT-1999; 99WO-US25561.

PR 30-OCT-1998; 98US-0106373.

PR 02-NOV-1998; 98US-0106537.

PA (INTE-) INTERLINK BIOTECHNOLOGIES LLC.

PA (KENT) UNIV KENTUCKY RES FOUND.

PI Everett NP, LI Q, Lawrence C, Davies MH;

DR WPI; 2000-365597/31.

PT Polypeptides for reducing proteolytic degradation of proteins administered to, or produced by a plant comprise indolicidin or its functional equivalents

PS Disclosure; Page 4; 50pp; English.

CC Indolicidin is a potent antimicrobial tridecapeptide, originally purified from cytoplasmic granules of bovine neutrophils. CP-11 is an analogue, which has better activity against *E. coli*, *Pseudomonas aeruginosa* and *Candida albicans*, but reduced activity against *Staphylococcus aureus*. A reverse peptide, Rev4 (AAY92796) of indolicidin was found to have increased stability against plant protease degradation. Expression of antimicrobial peptides in transgenic plants suffers a major limitation in that the foreign peptides are susceptible to rapid degradation by proteases. The invention concerns reducing the extent of protease degradation of a protein applied to, or produced by a plant by administering indolicidin, Rev4 or a functional equivalent to the plant. Transgenic plants expressing indolicidin and Rev4 are useful for production of the antimicrobial peptides. Compositions containing indolicidin and Rev4 are also useful for production of agronomically important proteins in plants.

CC Sequence 13 AA;

Query Match 100.0%; Score 91; DB 21; Length 13;
Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKWPMPWRK 13

DB 1 ILKWPMPWRK 13

RESULT 7
AAY91773
ID AAY91773 standard; Peptide; 13 AA.

XX

AC AAY91773;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment; leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma; breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon; multidrug resistance.

OS Synthetic.

PN WO9965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;

DR WPI; 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated polyoxalkylene-modified cationic peptides, useful for treating tumours

PS Disclosure; Page 14; 94pp; English.

CC This sequence represents a cationic peptide amino acid sequence, which can be used in the pharmaceutical composition of the invention. The invention relates to a pharmaceutical composition containing at least one activated polyoxalkylene (APO)-modified cationic peptide. The modification of peptides with APO increases their activity against tumour cells, including those with a multidrug resistant phenotype. The pharmaceutical composition can be used to treat tumours, specifically lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary, cervix, uterus, skin, prostate, liver and colon.

CC Sequence 13 AA;

Query Match 100.0%; Score 91; DB 21; Length 13;
Best Local Similarity 100.0%; Pred. No. 7.2e-07;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKWPMPWRK 13

DB 1 ILKWPMPWRK 13

RESULT 8
AAY91774
ID AAY91774 standard; Peptide; 13 AA.

AC AAY91774;

DT 06-JUN-2000 (first entry)

DE Amino acid sequence of cationic peptide MBI 11CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment; leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma; breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon; multidrug resistance.

OS Synthetic.

PN WO9965506-A2.

XX 23-DEC-1999.
 PD 14-JUN-1999; 99WO-CA00552.
 XX 12-JUN-1998; 98US-0096541.
 XX (MICR-) MICROLOGIX BIOTECH INC.
 PA Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP.
 PI WPI; 2000-223549/19.
 XX Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS
 XX Example 3; Page 14; 94pp; English.
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 13 AA;
 Query Match 100.0%; Score 91; DB 21; Length 13;
 Best Local Similarity 100.0%; Pred. No. 7.2e-07;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ILKKPMPMPMRK 13
 DB 1 ILKKPMPMPMRK 13
 RESULT 9
 ID AAY91818 standard; Peptide; 13 AA.
 XX AAY91818;
 AC 06-JUN-2000 (first entry).
 DT Amino acid sequence of cationic peptide MBI 11E1CN.
 DE
 XX Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 XX multidrug resistance.
 XX Synthetic.
 OS
 XX W09965506-A2.
 PN 23-DEC-1999.
 PD 14-JUN-1999; 99WO-CA00552.
 XX 12-JUN-1998; 98US-0096541.
 PR (MICR-) MICROLOGIX BIOTECH INC.
 XX Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 DR Novel pharmaceutical composition containing optionally activated
 XX

PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PT Disclosure; Page 15; 94pp; English.
 XX This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 XX
 SQ Sequence 13 AA;
 Query Match 100.0%; Score 91; DB 21; Length 13;
 Best Local Similarity 100.0%; Pred. No. 7.2e-07;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 ILKKPMPMPMRK 13
 DB 1 ILKKPMPMPMRK 13
 RESULT 10
 ID AAY91819 standard; Peptide; 13 AA.
 XX AAY91819;
 AC 06-JUN-2000 (first entry)
 DT Amino acid sequence of cationic peptide MBI 11E2CN.
 DE
 XX Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
 KW leukaemia; polyoxyalkylene-modified; APO; lymphoma; multiple myeloma;
 KM breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 XX multidrug resistance.
 XX Synthetic.
 OS
 XX W09965506-A2.
 PN 23-DEC-1999.
 PD 14-JUN-1999; 99WO-CA00552.
 XX 12-JUN-1998; 98US-0096541.
 PR (MICR-) MICROLOGIX BIOTECH INC.
 XX Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 PI WPI; 2000-223549/19.
 DR Novel pharmaceutical composition containing optionally activated
 PT polyoxyalkylene-modified cationic peptides, useful for treating tumours
 PS Disclosure; Page 15; 94pp; English.
 XX This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxyalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.

XX Sequence 13 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 21; Length 13;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWRRK 13
| | | | |
Db 1 ILKKPMPWRRK 13

RESULT 11

AA91820 ID AAY91820 standard; Peptide; 13 AA.

AC AAY91820;

DT 06-JUN-2000 (first entry);

DE Amino acid sequence of cationic peptide MBI 11E3CN.

KW Cationic peptide; tumour; pharmaceutical composition; cancer; treatment;
KW leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
KW multidrug resistance.

OS Synthetic.

PN WO965506-A2.

PD 23-DEC-1999.

PF 14-JUN-1999; 99WO-CA00552.

PR 12-JUN-1998; 98US-0096541.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
XX WPI: 2000-223549/19.

PT Novel pharmaceutical composition containing optionally activated
PT polyoxalkylene-modified cationic peptides, useful for treating tumours

PS Claim 1; Page 15; 94pp; English.

XX This sequence represents a cationic peptide amino acid sequence, which
CC can be used in the pharmaceutical composition of the invention. The
CC invention relates to a pharmaceutical composition containing at least one
CC activated polyoxalkylene (APO)-modified cationic peptide. The
CC modification of peptides with APO increases their activity against tumour
CC cells, including those with a multidrug resistant phenotype. The
CC pharmaceutical composition can be used to treat tumours, specifically
CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
CC cervix, uterus, skin, prostate, liver and colon.

XX Sequence 13 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 21; Length 13;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWRRK 13
| | | | |
Db 1 ILKKPMPWRRK 13

RESULT 12

AA924583 ID AAY24583 standard; peptide; 14 AA.

AC AAY24583;

DT 18-AUG-1999 (first entry)

DE Indolicidin analogue #35.

KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
KW antimicrobial; antibiotic; antitumour; surface disinfectant;
KW additive; shampoo; soap; insecticide; herbicide; preservative;
KW food; technical material.

OS Synthetic.

PN WO9807745-A2.

PD 26-FEB-1998.

PF 21-AUG-1997; 97WO-US14779.

PR 13-JAN-1997; 97US-0034949.

PR 21-AUG-1996; 96US-0024754.

PA (MICR-) MICROLOGIX BIOTECH INC.

PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MHP;
XX WPI: 1998-169090/15.

PT New indolicidin analogues with antimicrobial activity and related
PT nucleic acid - vectors, transformed cells and antibodies, also
PT conjugates with polyoxalkylene glycol and fatty acid to reduce
PT toxicity, useful therapeutically, as disinfectants etc.

PS Claim 13; Page 89; 129pp; English.

XX AAY24549 to AAY24615 represent indolicidin analogues of formulae
CC (I)-(VIII) containing up to 25 amino acids (aa): R₁XX₁XB₁(I), B₁XX₁XB₁
CC (II), B₁XX₁XB₁ (III), B₁XX₁XB₁B₁(AA)n₁LB₁B₁ (IV), B₁XX₁XB₁B₁(AA)n₁
CC (V), B₁XX₁XB₁XX₁XB₁ (VI), L₁XX₁XB₁ (VII) and B₁XX₁XB₁B₁ (VIII).
CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa,
CC preferably R or K; AA = any aa; n = 0 or 1; in (II), at least 1 Z = V;
CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
CC infections caused by bacteria (Gram positive or negative, or anaerobic);
CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
CC trematodes) or viruses. Typical of very many pathogens that can be
CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
CC aureus, Listeria, Clostridium, rotavirus and papilloma virus. Compounds
CC also derived from the analogues may be used similarly; the compounds may
CC may be used therapeutically or to coat medical devices; also they are
CC useful as surface disinfectants, as additives to shampoo or soaps, as
CC insecticides or herbicides, or as preservatives for foods and technical
CC materials. The analogues are administered by injection, lavage, orally
CC or topically, generally at 0.1-50 mg/kg. These analogues have a broader
CC spectrum of activity than indolicidin and modification as compounds
CC reduces their toxicity.

XX Sequence 14 AA;

Query Match
Best Local Similarity 100.0%; Score 91; DB 19; Length 14;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWRRK 13
| | | | |
Db 1 ILKKPMPWRRK 13

RESULT 13
 AAY91811
 ID AAY91811 standard; Peptide: 14 AA.
 XX
 AC AAY91811;
 XX
 DT 06-JUN-2000 (first entry)
 XX
 DE Amino acid sequence of cationic peptide MBI 11D11H.
 XX
 KW Cationic peptide: tumour; pharmaceutical composition: cancer; treatment;
 KW leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;
 KW multidrug resistance.
 XX
 OS Synthetic.
 XX
 PN W09965506-A2.
 XX
 PD 23-DEC-1999.
 XX
 PE 14-JUN-1999; 99WO-CA00552.
 XX
 PR 12-JUN-1998; 98US-0096541.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 XX
 DR WPI: 2000-223549/19.
 XX
 PT Novel pharmaceutical composition containing optionally activated
 PT polyoxalkylene-modified cationic peptides, useful for treating tumours
 PT
 XX
 PS Disclosure; Page 15; 94pp; English.
 XX
 SS This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 CC
 XX
 SO Sequence 14 AA:

Query Match 100.0%; Score 91; DB 21; Length 14;
 Best Local Similarity 100.0%; Pred. No. 7.8e-07;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRRK 13
 |||||
 DB 1 ILKKPMPWMPRRK 13

RESULT 14
 AAY24582
 ID AAY24582 standard; Peptide: 21 AA.
 XX
 AC AAY24582;
 XX
 DT 18-AUG-1999 (first entry)
 XX
 DE Indolicidin analogue #34.
 XX
 KW Indolicidin; bacterial infection; photo-oxidised solubiliser;
 KW antimicrobial; antidiabetic; antiarrhythmic; surface disinfectant;
 KW additive; shampoo; soap; insecticide; herbicide; preservative;
 KW food; technical material.

XX
 OS Synthetic.
 XX
 PN W09807745-A2.
 XX
 PD 26-FEB-1998.
 XX
 PE 21-AUG-1997; 97WO-US14779.
 XX
 PR 13-JAN-1997; 97US-0034949.
 XX
 PR 21-AUG-1996; 96US-0024754.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Erfle D, Fraser JR, Krieger TJ, Taylor R, West MH;
 XX
 DR WPI: 1998-169090/15.
 XX
 PT New indolicidin analogues with antimicrobial activity and related
 PT nucleic acid - vectors, transformed cells and antibodies, also
 PT conjugates with polyoxalkylene glycol and fatty acid to reduce
 PT toxicity, useful therapeutically, as disinfectants etc.
 XX
 PS Claim 13; Page 89; 129pp; English.
 XX
 SS AAY24549 to AAY24615 represent indolicidin analogues of formulae
 CC (I)-(VIII) containing up to 25 amino acids (aa): RXXXXZXB (I), BXXXXZXB
 CC (II), BXBXXZXB (III), BXXXXZXBn(A)nMILBBSGS (IV), BXXXXZXB(A)nM
 CC (V), LBBnXnXnXnXnXnX (VI), LKnXnXnXnXnXnX (VII) and BXXXXZXBn (VIII).
 CC Where Z = P or V; X = hydrophobic residue, preferably W; B = basic aa;
 CC preferably R or K; AA = any aa; n = 0 or 1; In (II), at least 1 Z = V;
 CC in (VIII) at least 2 X = F or Y. The analogues are used to treat
 CC infections caused by bacteria (Gram positive or negative, or anaerobic);
 CC fungi (yeast or moulds); parasites (protozoa, nematodes, cestodes or
 CC trematodes) or viruses. Typical of very many pathogens that can be
 CC controlled are Leishmania, Trypanosoma, Ascaris lumbricoides, Fasciola
 CC hepatica, Klebsiella pneumoniae, Bordetella pertussis, Staphylococcus
 CC aureus, listeria, Clostridium, rotavirus and papilloma virus. Compounds
 CC also be prepared from antibiotics or antiarrhythmic agents. The analogues
 CC may be used therapeutically or to coat medical devices; also they are
 CC useful as surface disinfectants, as additives to shampoo or soaps, as
 CC insecticides or herbicides, or as preservatives for foods and technical
 CC materials. The analogues are administered by injection, lavage, orally
 CC or topically, generally at 0.1-50 mg/Kg. These analogues have a broader
 CC spectrum of activity than indolicidin and modification as compounds
 CC reduces their toxicity.
 CC
 XX
 SO Sequence 21 AA:

Query Match 100.0%; Score 91; DB 19; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.2e-06;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRRK 13
 |||||
 DB 1 ILKKPMPWMPRRK 13

RESULT 15
 AAY91806
 ID AAY91806 standard; Peptide: 21 AA.
 XX
 AC AAY91806;
 XX
 DT 06-JUN-2000 (first entry)
 XX
 DE Amino acid sequence of cationic peptide MBI 11D4CN.
 XX
 KW Cationic peptide: tumour; pharmaceutical composition: cancer; treatment;
 KW leukaemia; polyoxalkylene-modified; APO; lymphoma; multiple myeloma;
 KW breast; lung; ovary; cervix; uterus; skin; prostate; liver; colon;

KM multidrug resistance.
 XX
 OS Synthetic.
 XX
 PN WO9965506-A2.
 XX
 PD 23-DEC-1999.
 XX
 PF 14-JUN-1999; 99WO-CA00552.
 XX
 PR 12-JUN-1998; 98US-0096541.
 XX
 PA (MICR-) MICROLOGIX BIOTECH INC.
 XX
 PI Friedland HD, Krieger TJ, Taylor R, Erfle D, Fraser JR, West MHP;
 XX
 DR WPI; 2000-223549/19.
 XX
 XX
 PT Novel pharmaceutical composition containing optionally activated
 PT polyoxaalkylene-modified cationic peptides, useful for treating tumours
 PT
 PS Disclosure; Page 15; 94pp; English.
 XX
 CC This sequence represents a cationic peptide amino acid sequence, which
 CC can be used in the pharmaceutical composition of the invention. The
 CC invention relates to a pharmaceutical composition containing at least one
 CC activated polyoxaalkylene (APO)-modified cationic peptide. The
 CC modification of peptides with APO increases their activity against tumour
 CC cells, including those with a multidrug resistant phenotype. The
 CC pharmaceutical composition can be used to treat tumours, specifically
 CC lymphoma, leukaemia, multiple myeloma, or tumours of breast, lung, ovary,
 CC cervix, uterus, skin, prostate, liver and colon.
 CC
 SQ Sequence 21 AA;

Query Match 100.0%; Score 91; DB 21; Length 21;
 Best Local Similarity 100.0%; Pred. No. 1.2e-06;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 ILKKPMPMPRRK 13
 |||||
 Db 1 ILKKPMPMPRRK 13

Search completed: January 4, 2002, 08:40:26
 Job time: 109 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:57 ; Search time 24.75 Seconds
(without alignments)
11.820 Million cell updates/sec

Title: US-09-444-281-35

Sequence: 1 ILKKWPMWPMRRK 13

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 212252 seqs, 22503292 residues

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	91	100.0	13	US-08-915-314-30	Sequence 30, Appl
2	91	100.0	13	US-08-915-314-62	Sequence 62, Appl
3	91	100.0	13	US-08-915-314-63	Sequence 63, Appl
4	91	100.0	13	US-08-915-314-64	Sequence 64, Appl
5	91	100.0	13	US-09-042-071-36	Sequence 36, Appl
6	91	100.0	14	US-08-915-314-57	Sequence 57, Appl
7	91	100.0	21	US-08-915-314-54	Sequence 54, Appl
8	87	95.6	12	US-08-915-314-52	Sequence 52, Appl
9	86	94.5	12	US-08-915-314-74	Sequence 74, Appl
10	86	94.5	12	US-08-702-054B-5	Sequence 5, Appl
11	86	94.5	13	US-08-915-314-58	Sequence 58, Appl
12	86	94.5	14	US-08-915-314-59	Sequence 59, Appl
13	86	94.5	15	US-08-702-054B-40	Sequence 40, Appl
14	86	94.5	20	US-08-915-314-55	Sequence 55, Appl
15	86	94.5	21	US-08-915-314-56	Sequence 56, Appl
16	85	93.4	12	US-08-915-314-38	Sequence 38, Appl
17	85	93.4	13	US-08-915-314-39	Sequence 39, Appl
18	85	93.4	13	US-08-915-314-45	Sequence 45, Appl
19	85	93.4	13	US-08-702-054B-1	Sequence 1, Appl
20	85	93.4	13	US-08-702-054B-17	Sequence 17, Appl
21	85	93.4	13	US-08-702-054B-32	Sequence 32, Appl
22	83	91.2	12	US-08-915-314-24	Sequence 24, Appl
23	83	91.2	13	US-08-915-314-49	Sequence 49, Appl
24	83	91.2	13	US-08-915-314-50	Sequence 50, Appl
25	83	91.2	13	US-08-915-314-51	Sequence 51, Appl
26	83	91.2	13	US-08-702-054B-30	Sequence 30, Appl
27	83	91.2	13	US-08-702-054B-31	Sequence 31, Appl

28	83	91.2	13	US-08-702-054B-34	Sequence 34, Appl
29	83	91.2	13	US-08-702-054B-35	Sequence 35, Appl
30	82	90.1	13	US-08-915-314-25	Sequence 25, Appl
31	82	90.1	13	US-08-915-314-66	Sequence 66, Appl
32	82	90.1	13	US-08-915-314-67	Sequence 67, Appl
33	82	90.1	13	US-08-702-054B-33	Sequence 33, Appl
34	81	89.0	11	US-08-915-314-75	Sequence 75, Appl
35	81	89.0	15	US-08-702-054B-39	Sequence 39, Appl
36	80	87.9	14	US-08-702-054B-18	Sequence 18, Appl
37	80	87.9	15	US-08-702-054B-41	Sequence 41, Appl
38	80	87.9	16	US-08-702-054B-2	Sequence 2, Appl
39	79.5	87.4	17	US-08-702-054B-38	Sequence 38, Appl
40	79	86.8	16	US-08-702-054B-42	Sequence 42, Appl
41	78	85.7	11	US-08-915-314-28	Sequence 28, Appl
42	78	85.7	12	US-08-915-314-40	Sequence 40, Appl
43	77	84.6	12	US-08-915-314-39	Sequence 39, Appl
44	77	84.6	12	US-08-702-054B-27	Sequence 27, Appl
45	76	83.5	12	US-08-915-314-77	Sequence 77, Appl

ALIGNMENTS

RESULT 1
US-08-915-314-30
Sequence 30, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
City: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-30

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ILKKWPMWPMRRK 13

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Db 1 ILKKPMPWPRRK 13
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RESULT 2
US-08-915-314-62.
; Sequence 62, Application US/08915314
; Patent No. 6180604
; GENERAL INFORMATION:
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erile, Douglas
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
; NUMBER OF SEQUENCES: 90
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/915,314
; FILING DATE: 20-AUG-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6180604tenburg Ph.D., Carol
; REGISTRATION NUMBER: 39,317
; REFERENCE/DOCKET NUMBER: 660081.405
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 62:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 13 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 1
; OTHER INFORMATION: /note="D-Form of Isoleucine"
US-08-915-314-62

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPRRK 13
DB 1 ILKKPMPWPRRK 13

RESULT 3
US-08-915-314-63
; Sequence 63, Application US/08915314
; Patent No. 6180604
; GENERAL INFORMATION:
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erile, Douglas
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
```

```
; TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
; NUMBER OF SEQUENCES: 90
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/915,314
; FILING DATE: 20-AUG-1997
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6180604tenburg Ph.D., Carol
; REGISTRATION NUMBER: 39,317
; REFERENCE/DOCKET NUMBER: 660081.405
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 63:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 13 amino acids
; TYPE: amino acid
; STRANDEDNESS:
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: Modified-site
; LOCATION: 13
; OTHER INFORMATION: /note="D-Form of Lysine"
US-08-915-314-63

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWPRRK 13
DB 1 ILKKPMPWPRRK 13

RESULT 4
US-08-915-314-64
; Sequence 64, Application US/08915314
; Patent No. 6180604
; GENERAL INFORMATION:
; APPLICANT: Fraser, Janet R.
; APPLICANT: West, Michael H.P.
; APPLICANT: Krieger, Timothy J.
; APPLICANT: Taylor, Robert
; APPLICANT: Erile, Douglas
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
; TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
; NUMBER OF SEQUENCES: 90
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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APPLICATION NUMBER: US/08/915:314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 64:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
FEATURE:
NAME/KEY: Modified-site
LOCATION: 1
OTHER INFORMATION: /note="D-Form of Isoleucine"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 13
OTHER INFORMATION: /note="D-Form of Lysine"
US-08-915-314-64

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPWRRK 13
Db 1 ILKKPMPWMPWRRK 13

RESULT 5
US-09-042-071-36
Sequence 36, Application US/09042071
Patent No. 6294372
GENERAL INFORMATION:
APPLICANT: Burian, Jan
APPLICANT: Kay, William W.
TITLE OF INVENTION: REPLICATION GENES AND GENE PRODUCTS FROM
TITLE OF INVENTION: SMALL CRYPTIC PLASMIDS AND METHODS FOR CONSTRUCTING
TITLE OF INVENTION: CONTROLLED-REPLICATION PLASMID VECTORS
NUMBER OF SEQUENCES: 52
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/042.071
FILING DATE: 13-MAR-1998
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 660081.407
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 13 amino acids

TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-042-071-36

Query Match 100.0%; Score 91; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 5.4e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPWRRK 13
Db 1 ILKKPMPWMPWRRK 13

RESULT 6
US-08-915-314-57
Sequence 57, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915.314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 57:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-57

Query Match 100.0%; Score 91; DB 4; Length 14;
Best Local Similarity 100.0%; Pred. No. 5.8e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPWRRK 13
Db 1 ILKKPMPWMPWRRK 13

RESULT 7
US-08-915-314-54
Sequence 54, Application US/08915314
Patent No. 6180604

GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESS: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 54:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-54

Query Match 100.0%; Score 91; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 8.5e-07;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPMPWPRRK 13
Db 1 ILKKPMPWPRRK 13

RESULT 8
US-08-915-314-52
Sequence 52, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESS: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 74:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 52:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-52

Query Match 95.6%; Score 87; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 1.8e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 LKKPMPWPRRK 13
Db 1 LKKPMPWPRRK 12

RESULT 9
US-08-915-314-74
Sequence 74, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESSES:
ADDRESS: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 74:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear

US-08-915-314-74

Query Match 94.5%; Score 86; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 2.4e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRR 12
Db 1 ILKKPMPWMPRR 12

RESULT 10

US-08-702-054B-5
Sequence 5, Application US/08702054B
Patent No. 6191254
GENERAL INFORMATION:
APPLICANT: Falls, Timothy J.
APPLICANT: Hancock, Robert E. W.
APPLICANT: Gough, Monisha
TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
TITLE OF INVENTION: AND METHODS OF SCREENING FOR THE SAME
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/702.054B
FILING DATE: 23-AUG-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/002.687
FILING DATE: 23-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Halle, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07420/013001
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 12 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-702-054B-5

Query Match 94.5%; Score 86; DB 4; Length 12;
Best Local Similarity 100.0%; Pred. No. 2.4e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRR 12
Db 1 ILKKPMPWMPRR 12

RESULT 11

US-08-915-314-58
Sequence 58, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.

APPLICANT: Krieger, Timothy J.

APPLICANT: Taylor, Robert

APPLICANT: Erfle, Douglas

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING

TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN

NUMBER OF SEQUENCES: 90

CORRESPONDENCE ADDRESS:

ADDRESSEE: SEED AND BERRY LLP

STREET: 6300 Columbia Center, 701 Fifth Avenue

CITY: Seattle

STATE: Washington

COUNTRY: USA

ZIP: 98104

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/915.314

FILING DATE: 20-AUG-1997

CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

NAME: No. 6180604tenburg Ph.D., Carol

REGISTRATION NUMBER: 39,317

REFERENCE/DOCKET NUMBER: 660081.405

TELEPHONE: (206) 622-4900

TELEFAX: (206) 682-6031

INFORMATION FOR SEQ ID NO: 58:

SEQUENCE CHARACTERISTICS:

LENGTH: 13 amino acids

TYPE: amino acid

STRANDEDNESS:

TOPOLOGY: linear

US-08-915-314-58

Query Match 94.5%; Score 86; DB 4; Length 13;
Best Local Similarity 100.0%; Pred. No. 2.0e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPMPWMPRR 12
Db 1 ILKKPMPWMPRR 12

RESULT 12

US-08-915-314-59
Sequence 59, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfle, Douglas
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED AND BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tendburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 59:
SEQUENCE CHARACTERISTICS:
LENGTH: 14 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-59

Query Match 94.5%; Score 86; DB 4; Length 14;
Best Local Similarity 100.0%; Pred. No. 2.8e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 13
US-08-702-054B-40
Sequence 40, Application US/08702054B
Patent No. 6191254
GENERAL INFORMATION:
APPLICANT: Palls, Timothy J.
APPLICANT: Hancock, Robert E. W.
TITLE OF INVENTION: ANTIMICROBIAL CATIONIC PEPTIDES
TITLE OF INVENTION: AND METHODS OF SCREENING FOR THE SAME
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSER: Fish & Richardson P.C.
STREET: 4225 Executive Square, Suite 1400
CITY: La Jolla
STATE: CA
COUNTRY: USA
ZIP: 92037
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/702,054B
FILING DATE: 23-AUG-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/002,687
FILING DATE: 23-AUG-1995
ATTORNEY/AGENT INFORMATION:
NAME: Haile, Lisa A.
REGISTRATION NUMBER: 38,347
REFERENCE/DOCKET NUMBER: 07420/013001
TELECOMMUNICATION INFORMATION:
TELEPHONE: 619/678-5070
TELEFAX: 619/678-5099
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 15 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-702-054B-40

Query Match 94.5%; Score 86; DB 4; Length 15;
Best Local Similarity 100.0%; Pred. No. 3e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 14
US-08-915-314-55
Sequence 55, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSER: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tendburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 55:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-55

Query Match 94.5%; Score 86; DB 4; Length 20;
Best Local Similarity 100.0%; Pred. No. 3.9e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ILKKPWPMPWR 12
DB 1 ILKKPWPMPWR 12

RESULT 15
US-08-915-314-56
Sequence 56, Application US/08915314
Patent No. 6180604
GENERAL INFORMATION:
APPLICANT: Fraser, Janet R.
APPLICANT: West, Michael H.P.
APPLICANT: Krieger, Timothy J.
APPLICANT: Taylor, Robert
APPLICANT: Erfile, Douglas

TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING
TITLE OF INVENTION: INFECTIONS USING ANALOGUES OF INDOLICIDIN
NUMBER OF SEQUENCES: 90
CORRESPONDENCE ADDRESS:
ADDRESSEE: SEED and BERRY LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: USA
ZIP: 98104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/915,314
FILING DATE: 20-AUG-1997
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: No. 6180604tenburg Ph.D., Carol
REGISTRATION NUMBER: 39,317
REFERENCE/DOCKET NUMBER: 660081.405
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 56:
SEQUENCE CHARACTERISTICS:
LENGTH: 21 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: linear
US-08-915-314-56

Query Match 94.58; Score 86; DB 4; Length 21;
Best Local Similarity 100.0%; Pred. No. 4.1e-06;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ILKKPWPWPWR 12
Db 1 ILKKPWPWPWR 12

Search completed: January 4, 2002, 08:40:57
Job time: 140 sec

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ORGANISM      Nicotiana benthamiana,
               Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
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               Asteridae; euasterids I; Solanales; Solanaceae; Nicotiana.
REFERENCE      1 (bases 1 to 6446)
AUTHORS        Gargier,S.J., Holtz,B.R., McCulloch,M.J. and Turner,T.H.
TITLE          A process for isolating and purifying viruses, soluble proteins and
               peptides from plant sources
JOURNAL        Patent: WO 011969-A 5 22-MAR-2001;
               Large Scale Biology Corporation (US)
FEATURES       source
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ORIGIN
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Percent Similarity: 100.000      Percent Identity: 100.000
alignment_block:
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Align seg 1/1 to: AX098418 from: 1 to: 6446
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seq_name: gb_htg:AP003754
seq_documentation_block:
LOCUS      AP003754 129052 bp DNA HTG 14-JUN-2001
DEFINITION Oryza sativa chromosome 7 clone OJ1341_A08, *** SEQUENCING IN
PROGRESS ***, in ordered pieces.
ACCESSION   AP003754
VERSION     AP003754.1 GI:14422472
KEYWORDS    HTG; HTGS_PHASE2.
SOURCE      Oryza sativa (cultivar:Nipponbare) DNA, clone:OJ1341_A08.
ORGANISM    Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
               Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
               Ehrhartoideae; Oryzaceae; Oryza.
REFERENCE    1 (sites)
AUTHORS      Sasaki,T., Matsumoto,T. and Yamamoto,K.
TITLE        Oryza sativa nipponbare(GA3) genomic DNA, chromosome 7, BAC
               clone:OJ1341_A08
JOURNAL      Published Only in DataBase (2001) In press
REFERENCE    2 (bases 1 to 129052)
AUTHORS      Sasaki,T., Matsumoto,T. and Yamamoto,K.
TITLE        Direct Submission
JOURNAL      Submitted (13-JUN-2001) Takuji Sasaki, National Institute of
               Agrobiological Resources, Rice Genome Research Program, Kannondai
               2-1-2, Tsukuba, Ibaraki 305-8602, Japan
               (E-mail:tsasaki@abrr.affrc.go.jp; URL:http://rgrp.dna.affrc.go.jp/,
               Tel:81-298-38-7441, Fax:81-298-38-7468)
COMMENT      The nucleotide sequence of this BAC clone was generated by
               combining Monsanto and RGP-Japan sequencing data.
               NOTE: It currently consists of 1 contigs. Gaps between the contigs
               are represented as runs of N. The order of the pieces is believed
               to be correct as given, however the sizes of the gaps between them
               are based on estimates that have provided by the submitter. This
               sequence will be replaced by the finished sequence as soon as it is
               available and the accession number will be preserved.
               * NOTE: This is a 'working draft' sequence.
               * This sequence will be replaced
               * by the finished sequence as soon as it is available and
               * the accession number will be preserved.
               Location/Qualifiers
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FEATURES       source

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/cultivar="Nipponbare"
/db_xref="taxon:4530"
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  Ratio: 6.800      Gaps: 0
Percent Similarity: 90.909      Percent Identity: 63.636
alignment_block:
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               :::::|||||||||||||||||
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seq_name: gb_ro:MM008210
seq_documentation_block:
LOCUS      MM008210 2651 bp mRNA ROD 31-OCT-1995
DEFINITION Mus musculus tropoelastin mRNA, complete cds.
ACCESSION   U08210
VERSION     U08210.1 GI:473273
KEYWORDS    house mouse.
SOURCE      Mus musculus
ORGANISM    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
               Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE    1 (bases 1 to 2651)
AUTHORS      Wyder,K.S., Sechler,J.L., Boyd,C.D. and Passmore,H.C.
TITLE        Use of an intron polymorphism to localize the tropoelastin gene to
               mouse chromosome 5 in a region of linkage conservation with human
               chromosome 7
JOURNAL      Genomics 23 (1), 125-131 (1994)
MEDLINE     95130069
REFERENCE    2 (bases 1 to 2651)
AUTHORS      Boyd,C.D.
TITLE        Direct Submission
JOURNAL      Submitted (30-MAR-1994) Charles D. Boyd, Department of Surgery,
               UMDNJ - Robert Wood Johnson Medical School, 51 French St., New
               Brunswick, NJ 08903, USA
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               GVGAGVGVGVGIGIGIGLSTGAVVPGAGIGAGKRGKRVGVLPGVPGVGVLP
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               VAVAVGAGVPGVAGTPAAAAAAKAAKAAKAGLGGVGGVGGVGIIPGVGVGAG
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    Ratio:        7.333         Gaps:        0
    Percent Similarity: 90.000   Percent Identity: 70.000

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US-09-444-281-35 x MMU08210 ..

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seq_name: gb_ro:AF289665

seq_documentation_block:
LOCUS       AF289665             107257 bp            DNA                ROD              14-AUG-2000
DEFINITION Mus musculus Eif4H gene, partial cds; LIMK1 gene, complete cds; and
ACCESSION   AF289665
VERSION     AF289665.1
KEYWORDS    GI:9800517

SOURCE      house mouse.
ORGANISM    Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE   1 (bases 1 to 107257)
AUTHORS    Green,E.D.
TITLE       Direct Submission
JOURNAL    Submitted (26-JUL-2000) Genome Technology Branch, National Human
Genome Research Institute, 49 Convent Dr. Rm. 2A02, Bethesda, MD
20892, USA

FEATURES             Location/Qualifiers
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                     33634..33720,34257..34440,37341..37507,37617..37722,
                     38745..38951,41098..41207,41439..41577,55071..55167,
                     56768..56882))
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     CDS              complement(join(25650..25812,25920..26077,26336..26391,
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GVDTLEINTEPIINRVPLDELDLTOETSLRLOTLHPDSDIGHGVSDSPVANSITH
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VGNCPYMAPEMINGRSYDEKVDYFESGTCVLCETICGNVADDTLPRMDGLVNRGFE
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P35=0.228, P36=0.969, P37=0.993, P38=0.905, P39=0.655,
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BASE COUNT 26300 a 26598 g 27966 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-09-444-281-35 x AF289665/rev
Align seg 1/1 to reverse of: AF289665 from: 1 to: 107257

4 LysTrpProTrpTrpProTrpArgArgLys 13

AGGTGGCCTTGTCGTCGAGCTCTCG 76468

seq_name: gb_htg:AC091250

seq_documentation_block:

LOCUS AC091250 200849 bp DNA HTG 11-APR-2001
DEFINITION Mus musculus chromosome 5 clone RP23-315E2 strain C57BL6/J, WORKING
DRAFT SEQUENCE, 7 unordered pieces.

AC091250

AC091250.1 GI:13592171

HTG: HTGS_PHASE1; HTGS_DRAFT.

house mouse.

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 200849)

Ayala, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W.,

Boutard, G.G., Brinkley, C., Brooks, S., Dietrich, N.L., Granito, S.,

Guan, X., Gupta, J., Ho, S.-L., Idol, J.R., Karlins, E., Lee-Lin, S.-O.,

Legaspi, R., Lim, M., Maduro, Q.L., Maduro, V.B., Masello, C.,

Masterian, S.D., McCloskey, J.C., McPowell, J., Pearson, R., Prasad, A.,

Shaychenko, Y., Snyder, B., Stantrop, S., Thomas, J.W., Thomas, P.J.,

Thompson, E.E., Touchman, J.W., Tsurgunov, C., Vogt, J.L., Walker, M.A.,

Welshy, R.D., Zhang, L.H. and Green, E.D.

NISC Comparative Sequencing Initiative

Unpublished

2 (bases 1 to 200849)

Green, E.D.

Direct Submission

Submitted (11-APR-2001) NIH Intramural Sequencing Center, 8717

Groveport Circle, Gaithersburg, MD 20877, USA

Genome Center

Center: NIH Intramural Sequencing Center

Center code: NISC

Web site: http://www.nisc.nih.gov

Contact: nisc.mouse@nhgri.nih.gov

Project Information

Center project name: aty

Center clone name: 315E02

Summary Statistics

Sequencing vector: plasmid; n/a; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap, version 0.990319

Consensus quality: 197727 bases at least Q40

Consensus quality: 198423 bases at least Q30

Consensus quality: 198823 bases at least Q20

Insert size: 202000; agarose-fp

Insert size: 200249; sum-of-contigs

Quality coverage: 9.87x in Q20 bases; sum-of-contigs

Quality coverage: 9.95x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently

consists of 7 contigs. The true order of the pieces

is not known and their order in this sequence record is

arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

as soon as it is available and the accession number will

be preserved.

1 2812: contig of 2812 bp in length

2813 2812: gap of unknown length

2913 9301: contig of 6389 bp in length

9302 9401: gap of unknown length

9402 17173: contig of 7772 bp in length

17174 17273: gap of unknown length

17274 26044: contig of 8771 bp in length

26045 26144: gap of unknown length

FEATURES
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/organism="Mus musculus"
/strain="C57BL6/J"
/db_xref="taxon:10090"
/chromosome="5"
/clone="RP23-315E2"
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9402 17173
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17274 26044
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26145 65929
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clone_end:SP6
vector_side:right"
66030 109036
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vector_side:right"

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65930 66029: gap of unknown length
66030 109036: contig of 43007 bp in length
109037 109136: gap of unknown length
109137 200849: contig of 91713 bp in length.
Location/Qualifiers

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-09-444-281-35 x AC091250 ..

Align seg 1/1 to: AC091250 from: 1 to: 200849

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AGGTGGCCTTGTCGTCGAGCTCTCG 148482

seq_name: gb_htg:AC020365

seq_documentation_block:

LOCUS AC020365 27780 bp DNA HTG 03-JAN-2000

DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS *** in ordered

pieces.

AC020365

AC020365.1 GI:6664532

HTG: HTGS_PHASE2.

fruit fly

Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;

Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;

Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

1 (bases 1 to 27780)

Adams, M. and Venter, J.C.

Direct Submission

Submitted (30-DEC-1999) Celera Genomics, 45 West Gude Drive,

Rockville, MD, USA

This sequence was identified as CDL:10212917 by the submitter.

For more information on this record e-mail to fly@celera.com.

NOTE: This is a 'working draft' sequence.

* This sequence will be replaced

* by the finished sequence as soon as it is available and

* the accession number will be preserved.

FEATURES
source 1. 2780
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"

BASE COUNT 7869 a 6128 c 6190 g 7593 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x AC020365 ..

Align seg 1/1 to: AC020365 from: 1 to: 27780

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seq_name: gb_in:AC008316

seq_documentation_block:

LOCUS AC008316 160817 bp DNA INV 06-MAR-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 85F-85F, BAC clone
BACR23004, complete sequence.

ACCESSION AC008316
VERSION AC008316.3 GI:13236627
KEYWORDS HTG.

SOURCE
ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 160817).
Celinker,S.E., Adams,M.D., Krommler,B., Tyler,D., Wan,K.H.,
Holt,R.A., Evans,C.A., Gocayne,J.D., Amanatides,P.G., Brandon,R.C.,
Rogers,Y., An,H., Baldwin,D., Banzon,J., Beeson,K.Y., Busam,D.A.,
Carlson,J.W., Center,A., Chape,M., Chavez,C., Chew,M., Ciesiolka,L.,
Dodson,K., Dorselt,V., Doup,L.E., Doyle,C., Dresnek,D., Farfan,D.,
Fierlier,S., Frise,E., Galle,R.F., Gary,N.S., George,R.A.,
Gonzalez,M., Houck,J., Hoskins,R.A., Hostin,D., Howland,T.J.,
Ibegam,C., Jatali,M., Kruse,D., Li,P., Mattei,B., Moshrefi,A.,
McIntosh,T.C., Moy,M., Murphy,B., Nelson,C., Nelson,K.A., Nunoo,J.,
Pacleb,J., Paragas,V., Park,S., Patel,S., Pfeiffer,B.,
Phouanavong,S., Pittman,G.S., Puri,V., Richards,S., Scheeler,F.,
Stapleton,M., Strong,R., Svirskas,R., Tector,C., Williams,S.M.,
Zaveri,J.S., Smith,H.O., Rubin,G.M. and Venter,J.C.

JOURNAL
TITLE
REFERENCE
AUTHORS
2 (bases 1 to 160817)
Celinker,S.E., Agapayni,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Chape,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karia,K., Kearney,L.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.
Direct Submission
Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Mar 6, 2001 this sequence version replaced gi:6957904.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.

For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (<http://www.fruitfly.org/sequence/>) or send email
to bdg@fruitfly.berkeley.edu.

FEATURES
source 1. 160817
/organism="Drosophila melanogaster"
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Drosophila melanogaster BAC library, partial EcoRI in
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ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x AC008316 ..

Align seg 1/1 to: AC008316 from: 1 to: 160817

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seq_name: gb_in:AC008315

seq_documentation_block:

LOCUS AC008315 177028 bp DNA INV 17-FEB-2001
DEFINITION Drosophila melanogaster, chromosome 3R, region 85E-85E, BAC clone
BACR23001, complete sequence.

ACCESSION AC008315
VERSION AC008315.9 GI:12957660
KEYWORDS HTG.

SOURCE
ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 177028).
Celinker,S.E., Adams,M.D., Krommler,B., Tyler,D., Wan,K.H.,
Holt,R.A., Evans,C.A., Gocayne,J.D., Amanatides,P.G., Brandon,R.C.,
Rogers,Y., An,H., Baldwin,D., Banzon,J., Beeson,K.Y., Busam,D.A.,
Carlson,J.W., Center,A., Chape,M., Davenport,L.B., Dietz,S.M.,
Dodson,K., Dorselt,V., Doup,L.E., Doyle,C., Dresnek,D., Farfan,D.,
Fierlier,S., Frise,E., Galle,R.F., Gary,N.S., George,R.A.,
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Ibegam,C., Jatali,M., Kruse,D., Li,P., Mattei,B., Moshrefi,A.,
McIntosh,T.C., Moy,M., Murphy,B., Nelson,C., Nelson,K.A., Nunoo,J.,
Pacleb,J., Paragas,V., Park,S., Patel,S., Pfeiffer,B.,
Phouanavong,S., Pittman,G.S., Puri,V., Richards,S., Scheeler,F.,
Stapleton,M., Strong,R., Svirskas,R., Tector,C., Williams,S.M.,
Zaveri,J.S., Smith,H.O., Rubin,G.M. and Venter,J.C.

JOURNAL
TITLE
REFERENCE
AUTHORS
2 (bases 1 to 177028)
Celinker,S.E., Agapayni,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Chape,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karia,K., Kearney,L.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.
Direct Submission
Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Mar 6, 2001 this sequence version replaced gi:6957904.
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
This sequence was assembled using end sequences from a whole genome
shotgun and from subclones of this BAC and its neighboring clones.


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alignment_scores:

Quality:	65.00	Length:	15
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Percent Similarity:	73.333	Percent Identity:	60.000

alignment_block:

US-09-444-281-35 x AE003684 ..

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seq_documentation_block:

LOCUS	MM421XCOA	19479 bp	DNA	08-APR-1994
DEFINITION	M.musculus alpha2 (IX) collagen gene, complete CDS.			
ACCESSION	222923			
VERSION	222923.1 GI:311949			
KEYWORDS	alpha2 (IX) collagen.			
SOURCE	house mouse.			
ORGANISM	Mus musculus			

REFERENCE

1 (bases 1 to 19479)
Perala,M., Elina,K., Metsaranta,M., Rosati,R., de Crombrughe,B.
and Vuorio,E.
The exon structure of the mouse alpha 2(IX) collagen gene shows
unexpected divergence from the mouse alpha 2(IX) collagen gene

AUTHORS

JOURNAL
MEDLINE
REFERENCE
94148964
2 (bases 1 to 19479)
Perala,M.
Direct Submission
Submitted (07-JUN-1993) Merja Perala, Medical Biochemistry,
University of Turku, Kilnamylykatu 10, 20520 Turku, Finland
Location/Qualifiers
1. 19479
/organism="Mus musculus"
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TITILE

JOURNAL
MEDLINE
REFERENCE
94148964
2 (bases 1 to 19479)
Perala,M.
Direct Submission
Submitted (07-JUN-1993) Merja Perala, Medical Biochemistry,
University of Turku, Kilnamylykatu 10, 20520 Turku, Finland
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TITILE

JOURNAL
MEDLINE
REFERENCE
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2 (bases 1 to 19479)
Perala,M.
Direct Submission
Submitted (07-JUN-1993) Merja Perala, Medical Biochemistry,
University of Turku, Kilnamylykatu 10, 20520 Turku, Finland
Location/Qualifiers
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FEATURES

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Align seg 1/1 to reverse of: MMA2IXCOA from: 1 to: 19479

5 TrpProTrpTrpProTrpargArgLys 13
|||||
4355 TGGCCCTGCTGGCCCTGAGGACCCGG 4329

seq_name: gb_pr:AL162423

```
seq_documentation_block:
```

DEFINITION Human DNA sequence from clone RP11-456D21 on chromosome 9, complete

sequence.

VERSION ALL62423.18 GI:13234851

KEYWORDS HTG.

ORGANISM

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE	1 (bases 1 to 88323)
ALTHOFFS	Carroll D

TITLE

COMMENT

where difference sequences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em., EMBL; Sw., SWISSPROT; Tr., TrEMBL; Wp., WormPep; Information on the WormPep database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr9> Rpl1-45621 is from the library RPlC-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm> VECTOR: pBAC3.6

IMPORTANT: This sequence is not the entire insert of clone Rpl1-45621. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone Rpl1-45621 is at 1 in this sequence. The true left end of clone Rpl1-49882 is at 88224 in this sequence. The true right end of clone Rpl1-297L11 is at 34844 in this sequence.

FEATURES	Location/Qualifiers
source	1. . 88323

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="9"
/clone="RP11-456D21"
/clone_id="RP11-11.2"
362. .588
/clone="HAL1 repeat: matches 595. .829 of consensus"
963. .1009
/clone="LMNC repeat: matches 2085. .2405 of consensus"
1168. .1751
/clone="L2 repeat: matches 2630. .3230 of consensus"
1175. .2047
/clone="L2 repeat: matches 2248. .2526 of consensus"
2061. .2215
/clone="MIR repeat: matches 82. .252 of consensus"
2443. .2788
/clone="L1M4 repeat: matches 5399. .5754 of consensus"
2889. .3184
/clone="L1 repeat: matches 1. .298 of consensus"
3306. .3446
/clone="FLM_C repeat: matches 2. .133 of consensus"
3862. .4181
/clone="ALUdJ repeat: matches 2. .312 of consensus"
5285. .5366
/clone="41 copies 2 mer aa 638 conserved"
5681. .5953
/clone="ALUSX repeat: matches 1. .395 of consensus"
6211. .6508
/clone="ALUdJ repeat: matches 2. .298 of consensus"
6566. .6961
/clone="MLT1 repeat: matches 3. .402 of consensus"
6989. .7284
/clone="ALUSg repeat: matches 1. .296 of consensus"
8131. .8238
/clone="MIR repeat: matches 124. .256 of consensus"
8437. .8740
/clone="ALUS repeat: matches 1. .304 of consensus"
8936. .9019
/clone="L2 repeat: matches 2660. .2749 of consensus"
9150. .9379
/clone="MIR repeat: matches 24. .259 of consensus"
10513. .10672
/clone="MIR repeat: matches 63. .218 of consensus"
10674. .10795
/clone="ALUSg/x repeat: matches 172. .293 of consensus"
11633. .11896
/clone="L2 repeat: matches 2414. .2682 of consensus"
11870. .12134
/clone="MIR repeat: matches 8. .261 of consensus"
12183. .12502

```


----- Project Information
Center project name: ba180A14
----- Summary Statistics
Assembly program: XGAP4; version 4.5
Sequencing vector: M13; M7815; 43% of reads
Sequencing vector: plasmid; L08752; 56% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 33302 bases at least Q40
Consensus quality: 341682 bases at least Q30
Consensus quality: 346727 bases at least Q20
Insert size: 351326; sum-of-contigs
Insert size: 176896; 2.8% error; agarose-fp
Quality coverage: 3.95x in Q20 bases; sum-of-contigs Quality
coverage: 8.06x in Q20 bases; agarose-fp

Draft Sequence Produced by Whitehead Institute/MIT Center for
Genome Research, 320 Charles Street,
Cambridge, MA 02141, USA
http://www-seq.wi.mit.edu.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 38 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 16446: contig of 16446 bp in length
16447 16546: gap of 100 bp
16547 22746: contig of 6200 bp in length
22747 22846: gap of 100 bp
22847 45482: contig of 22636 bp in length
45483 45582: gap of 100 bp
45583 49251: contig of 3663 bp in length
49252 49351: gap of 100 bp
49352 53239: contig of 3888 bp in length
53240 53339: gap of 100 bp
53340 58178: contig of 4839 bp in length
58179 58278: gap of 100 bp
58279 71754: contig of 13476 bp in length
71755 71854: gap of 100 bp
71855 76641: contig of 4787 bp in length
76642 76741: gap of 100 bp
76742 80415: contig of 3674 bp in length
80416 80515: gap of 100 bp
80516 89902: contig of 9387 bp in length
89903 90002: gap of 100 bp
90003 93661: contig of 3659 bp in length
93662 93761: gap of 100 bp
93762 101785: contig of 8024 bp in length
101786 101885: gap of 100 bp
101886 109333: contig of 7448 bp in length
109334 109433: gap of 100 bp
109434 121288: contig of 11855 bp in length
121289 121388: gap of 100 bp
121389 128767: contig of 7379 bp in length
128768 128867: gap of 100 bp
128868 131917: contig of 3050 bp in length
131918 132017: gap of 100 bp
132018 138795: contig of 6778 bp in length
138796 138895: gap of 100 bp
138896 144660: contig of 5765 bp in length
144661 144760: gap of 100 bp
144761 156587: contig of 11827 bp in length
156588 156687: gap of 100 bp
156688 158939: contig of 2252 bp in length
158940 159039: gap of 100 bp
159040 161353: contig of 2314 bp in length
161354 161453: gap of 100 bp
161454 165170: contig of 3717 bp in length
165171 165270: gap of 100 bp
165271 168479: contig of 3209 bp in length
168480 168579: gap of 100 bp

168580 172188: contig of 3609 bp in length
172189 172288: gap of 100 bp
172289 177084: contig of 4796 bp in length
177085 177184: gap of 100 bp
177185 199876: contig of 22692 bp in length
199877 199976: gap of 100 bp
199977 222186: contig of 22210 bp in length
222187 222286: gap of 100 bp
222287 240235: contig of 17949 bp in length
240236 240335: gap of 100 bp
240336 263715: contig of 23360 bp in length
263716 263815: gap of 100 bp
263816 269270: contig of 5455 bp in length
269271 269370: gap of 100 bp
269371 271745: contig of 2375 bp in length
271746 271845: gap of 100 bp
271846 277024: contig of 5179 bp in length
277025 277124: gap of 100 bp
277125 280037: contig of 2913 bp in length
280038 280137: gap of 100 bp
280138 307328: contig of 27191 bp in length
307329 307428: gap of 100 bp
307429 313985: contig of 6557 bp in length
313986 314085: gap of 100 bp
314086 330085: contig of 16000 bp in length
330086 330185: gap of 100 bp
330186 334887: contig of 4702 bp in length
334888 334987: gap of 100 bp
334988 355026: contig of 20039 bp in length.

FEATURES
source
1. 355026
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-180A14"
/clone_lib="RPC1-11.1"
1. 16446
/note="assembly_fragment:02644
fragment_chain:1"
16547. 22746
/note="assembly_fragment:02589
fragment_chain:1"
22847. 45482
/note="assembly_fragment:03035
fragment_chain:1"
45583. 49251
/note="assembly_fragment:00032"
49352. 53239
/note="assembly_fragment:00060"
53340. 58178
/note="assembly_fragment:00105"
58279. 71754
/note="assembly_fragment:00250"
71855. 76641
/note="assembly_fragment:00381"
76742. 80415
/note="assembly_fragment:00389"
80516. 89902
/note="assembly_fragment:00591"
90003. 93661
/note="assembly_fragment:00750"
93762. 101785
/note="assembly_fragment:00754"
101886. 109333
/note="assembly_fragment:00819"
109434. 121288
/note="assembly_fragment:00889"
121389. 128767
/note="assembly_fragment:00932"
128868. 131917
/note="assembly_fragment:00972"
132018. 138795
/note="assembly_fragment:01033"


```

* 91004 91103: gap of 100 bp
* 91104 109428: contig of 18325 bp in length
* 109429 109528: gap of 100 bp
* 109529 135277: contig of 25749 bp in length
* 135278 135377: gap of 100 bp
* 135378 159391: contig of 24014 bp in length.
Location/Qualifiers
1. 159391

```

FEATURES

source

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/map="5"

```

```

/clone="RP11-67212"
/clone.lib="RPC1-11 Human Male BAC"

```

misc_feature

1. 1373

misc_feature

1474. 3371

misc_feature

3472. 5991

misc_feature

6092. 8001

misc_feature

8102. 10452

misc_feature

10553. 13499

misc_feature

13600. 17300

misc_feature

17401. 23167

misc_feature

23268. 27394

misc_feature

clone_end:T7

misc_feature

vector_side:right

misc_feature

27495. 31941

misc_feature

32042. 36867

misc_feature

36968. 43237

misc_feature

43338. 52795

misc_feature

52896. 61446

misc_feature

61547. 68822

misc_feature

68923. 79907

misc_feature

80008. 91003

misc_feature

91104. 109428

misc_feature

109529. 135277

misc_feature

135378. 159391

misc_feature

clone_end:SP6

misc_feature

vector_side:right

misc_feature

35737 c 36282 g 41744 t 1901 others

misc_feature

BASE COUNT

ORIGIN

alignment_scores:

Quality: 64.00

Ratio: 7.111 Length: 9

Percent similarity: 100.000 Gaps: 0

Percent identity: 77.778

alignment_block:

US-09-444-281-35 x AC027113

Align seg 1/1 to: AC027113 from: 1 to: 159391

5 TrpProTPrpProTPrpAtrGly 13

87682 TGGCCGTGGCTTGGAGGAGGAG 87708

seq_name: gb_hgt:AL358473

seq_documentation_block:

LOCUS

AL358473 187845 bp DNA

Homo sapiens chromosome 1 clone RP11-168016, *** SEQUENCING IN

DEFINITION

Homo sapiens chromosome 1 clone RP11-168016, *** SEQUENCING IN

ACCESSION

AL358473 AC027712

VERSION

AL358473.8 GI:11322009

KEYWORDS

HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.

SOURCE

human.

ORGANISM

Homo sapiens

REFERENCE

McLay, K. Submitted (06-APR-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk

AUTHORS

On May 15, 2001 this sequence version replaced gi:7712176

TITLE

gi:10800605.

JOURNAL

Genome Center

COMMENT

Center: Sanger Centre

Center code: SC

Web site: http://www.sanger.ac.uk

Contact: humquerry@sanger.ac.uk

Project Information

Center project name: BA168016

Summary Statistics

Assembly program: XGAP4; version 4.5

Sequencing vector: M13; M77815; 56% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Consensus quality: 186132 bases at least Q40

Consensus quality: 186498 bases at least Q30

Consensus quality: 186851 bases at least Q20

Insert size: 187245; sum-of-contigs

Insert size: 186146; agarose-fp

Quality coverage: 10.30x in Q20 bases; sum-of-contigs Quality

coverage: 10.58x in Q20 bases; agarose-fp

Draft Sequence Produced by Whitehead Institute/MIT Center for

Genome Research, 320 Charles Street,

Cambridge, MA 02141, USA

http://www-seq.wi.mit.edu.

NOTE: This is a 'working draft' sequence. It currently

consists of 7 contigs. The true order of the pieces

is not known and their order in this sequence record is

arbitrary. Gaps between the contigs are represented as

runs of N, but the exact sizes of the gaps are unknown.

This record will be updated with the finished sequence

as soon as it is available and the accession number will

be preserved.

1 11961: contig of 11961 bp in length

11962 12061: gap of 100 bp

12062 95058: contig of 82997 bp in length

95059 95158: gap of 100 bp

95159 124623: contig of 29465 bp in length

124624 124723: gap of 100 bp

124724 128609: contig of 4886 bp in length

128610 129709: gap of 100 bp

129710 151381: contig of 21672 bp in length

151382 151481: gap of 100 bp

151482 174927: contig of 23446 bp in length

174928 175027: gap of 100 bp

175028 187845: contig of 12818 bp in length.

Location/Qualifiers

1. 187845

/organism="Homo sapiens"

/db_xref="taxon:9606"

```

misc_feature /chromosome="1"
/clone="Rp11-168016"
/clone_lib="RPC1-11.1"
1..11961
/note="assembly_fragment:01682
fragment_chain:1
clone_end:SP6
vector_side:left"
misc_feature 12062..95058
/note="assembly_fragment:01450
fragment_chain:1"
misc_feature 95159..124623
/note="assembly_fragment:01175
fragment_chain:1"
misc_feature 124724..129609
/note="assembly_fragment:04318
fragment_chain:1"
misc_feature 129710..151381
/note="assembly_fragment:00767
fragment_chain:2"
misc_feature 151482..174927
/note="assembly_fragment:01178
fragment_chain:2"
misc_feature 175028..187845
/note="assembly_fragment:02771
clone_end:R7
vector_side:right"
BASE COUNT 46168 a 48356 c 47958 g 44763 t 600 others
ORIGIN

```

```

alignment_scores:      Quality: 64.00      Length: 9
Ratio: 7.111           Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

```

alignment_block:
US-09-444-281-35 x AL358473

Align seg 1/1 to: AL358473 from: 1 to: 187845

```

5 TrpProTrrPrpTrrPrpAtrGlys 13
|||||
133213 TGGCCCTGTGGCTTGGAGGAGGAG 133239

```


[illegible]

```

XX 20-SEP-2000 (first entry)
DT
XX Oligonucleotide used for synthesis of MBI 2X11B7 poly cassette.
DE
XX Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
KM MBI-11; indolicidin; bovine; ss.
XX
XX Synthetic.
OS
XX WO200031279-A2.
PN
XX 02-JUN-2000.
PD
XX 19-NOV-1999; 99WO-CA01107.
PF
XX 20-NOV-1998; 98US-0109218.
PR
XX (MICR-) MICROLOGIX BIOTECH INC.
PA
XX
XX Burian J, Bartfeld D:
PI
XX WPI: 2000-400086/34.
DR
XX
XX Multi-domain fusion protein expression cassette used for high yield
PT stable production of foreign peptide gene products -
XX
XX Example 5; Page 39; 73pp: English.
PS
XX A novel method allows the efficient production of cationic peptides in
CC recombinant host cells. The method involves construction of a
CC multi-domain fusion protein expression cassette comprising a promoter and
CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
CC of anionic peptide sequences in the linker sequences neutralises the
CC positive charge of the cationic peptide so that the charge of the
CC fusion protein is controlled. This cassette allows high yield, stable
CC production of the cationic peptide. Cationic peptides such as
CC bovine indolicidin may be used as antimicrobial agents. The present
CC sequence is an oligonucleotide that was used to synthesise a
CC MBI-11B7 fragment. This fragment was used in the expression
CC cassette. MBI-11B7 is a cationic peptide derived from modifications
CC of indolicidin.
XX
XX Sequence 108 BP; 18 A; 33 C; 31 G; 26 T; 0 other;
SQ
XX
XX Alignment_scores:
XX Quality: 76.00 Length: 13
XX Ratio: 6.333 Gaps: 0
XX Percent Similarity: 92.308 Percent Identity: 69.231
XX
XX alignment_block:
XX US-09-444-281-35 x AAA27296 ..
XX
XX Align seg 1/1 to: AAA27296 from: 1 to: 108
XX
XX 1 lIleuLysLysTrpProTrrpTrpProTrrpArgArgLys 13
XX : : : : : : : : : : : : : : : : : : : : :
XX 38 AFGATCTGCGGTTGGCCGTGTGTGCGCGTGGCGCAAA 76
XX
XX seq_name: /SID58/gc9data/geneseq/geneseqn/NA2000.DAT:AAA27298
XX
XX seq_documentation_block:
XX ID AAA27298 standard; DNA; 114 BP.
XX
XX AAA27298;
XX
XX 20-SEP-2000 (first entry)
XX
XX Oligonucleotide used for synthesis of MBI 11B7 first cassette.
DE
XX Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
KM MBI-11; indolicidin; bovine; ss.
XX

```

```
XX OS Synthetic.
XX PN WO200031279-A2.
XX PD 02-JUN-2000.
XX PF 19-NOV-1999; 99WO-CA01107.
XX PR 20-NOV-1998; 98US-0109218.
XX PA (MICR-) MICROLOGIX BIOTECH INC.
XX PI Burlan J, Bartfeld D;
XX DR WPI; 2000-400086/34.
XX PT Multi-domain fusion protein expression cassette used for high yield
XX PS stable production of foreign peptide gene products -
XX PS Example 5; Page 40; 73pp; English.
XX CC A novel method allows the efficient production of cationic peptides in
XX CC recombinant host cells. The method involves construction of a
XX CC multi-domain fusion protein expression cassette comprising a promoter and
XX CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
XX CC of anionic peptide sequences in the linker sequences neutralises the
XX CC positive charge of the cationic peptide so that the charge of the
XX CC fusion protein is controlled. This cassette allows high yield, stable
XX CC production of the cationic peptide. Cationic peptides such as
XX CC bovine indolicidin may be used as antimicrobial agents. The present
XX CC sequence is an oligonucleotide that was used to synthesise a
XX CC MBI-11B7 fragment. This fragment was used in the expression
XX CC cassette. MBI-11B7 is a cationic peptide derived from modifications
XX CC of indolicidin.
SQ Sequence 114 BP; 20 A; 34 C; 32 G; 28 T; 0 other;

alignment_scores:
Quality: 76.00 Length: 13
Ratio: 6.333 Gaps: 0
Percent Similarity: 92.308 Percent Identity: 69.231

alignment_block:
US-09-444-281-35 x AAA27298
Align seg 1/1 to: AAA27298 from: 1 to: 114
1 ILEuLYSLySTRpTROTrPRpRrPARqARGLyS 13
++++: ::::::::::::::::::::::::::::
44 ATGATTCGCGTGTGGCCGTGGCGCGTGCACAA 82

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA27294
seq_documentation_block:
ID AAA27294 standard; DNA; 51 BP.
XX AC AAA27294;
XX DT 20-SEP-2000 (first entry)
XX DE Oligonucleotide used for synthesis of MBI 2X11B7 last cassette.
XX KW Oligonucleotide; cellulose binding domain; CBD; cationic peptide;
XX MBI-11; indolicidin; bovine; ss.
XX OS Synthetic.
XX PN WO200031279-A2.
XX PD 02-JUN-2000.
XX PF
```

```
PF 19-NOV-1999; 99WO-CA01107.
XX PR 20-NOV-1998; 98US-0109218.
XX PA (MICR-) MICROLOGIX BIOTECH INC.
XX PI Burlan J, Bartfeld D;
XX DR WPI; 2000-400086/34.
XX PT Multi-domain fusion protein expression cassette used for high yield
XX PS stable production of foreign peptide gene products -
XX PS Example 5; Page 38; 73pp; English.
XX CC A novel method allows the efficient production of cationic peptides in
XX CC recombinant host cells. The method involves construction of a
XX CC multi-domain fusion protein expression cassette comprising a promoter and
XX CC a nucleic acid molecule expressed as an insoluble protein. The inclusion
XX CC of anionic peptide sequences in the linker sequences neutralises the
XX CC positive charge of the cationic peptide so that the charge of the
XX CC fusion protein is controlled. This cassette allows high yield, stable
XX CC production of the cationic peptide. Cationic peptides such as
XX CC bovine indolicidin may be used as antimicrobial agents. The present
XX CC sequence is an oligonucleotide that was used to synthesise a
XX CC MBI-11B7 fragment. This fragment was used in the expression
XX CC cassette. MBI-11B7 is a cationic peptide derived from modifications
XX CC of indolicidin.
SQ Sequence 151 BP; 22 A; 44 C; 49 G; 36 T; 0 other;

alignment_scores:
Quality: 76.00 Length: 13
Ratio: 6.333 Gaps: 0
Percent Similarity: 92.308 Percent Identity: 69.231

alignment_block:
US-09-444-281-35 x AAA27294
Align seg 1/1 to: AAA27294 from: 1 to: 151
1 ILEuLYSLySTRpTROTrPRpRrPARqARGLyS 13
++++: ::::::::::::::::::::::::::::::
38 ATGATTCGCGTGTGGCCGTGGCGCGTGCACAA 76

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA1999.DAT:AAV83788
seq_documentation_block:
ID AAV83788 standard; DNA; 39 BP.
XX AC AAV83788;
XX DT 19-MAR-1999 (first entry)
XX DE Antimicrobial peptide Indolicidin encoding DNA.
XX KW Antimicrobial; fusion; acidic peptide; recombinant; microorganism;
XX guamerin; basic peptide; indolicidin; ss.
XX OS Synthetic.
XX OS Bos sp.
XX FH Key Location/Qualifiers
XX FT CDS 1..39
XX FT /*tag= a
XX FT /note= "the start and stop codons are not indicated"
XX PN WO9854336-A1.
XX PD 03-DEC-1998.
XX PF 28-MAY-1998; 98WO-KR00132.
```

```

XX 09-APR-1998; 98KR-0013372.
PR 28-MAY-1997; 97KR-0021312.
XX
XX (KOAD ) KOREA ADV INST SCI & TECHNOLOGY.
PA (SAMY-) SAMYANG GENEX CORP.
XX
PI Hong S, Kang MH, Kim JH, Kim S, Lee H, Lee JH;
XX WPI, 1999-059844/05.
DR P-PSDB; AAW87609.
XX
PT New method for mass production of antimicrobial peptides - by
PT constructing fusion genes, comprising acidic and antimicrobial
PT peptide genes and transforming host with vector containing these
XX
PS Example 6; Page 18; 52pp; English.
XX
CC The invention relates to mass production of antimicrobial peptides. The
CC method comprises constructing a fusion gene containing a first gene
CC encoding a negatively charged acidic peptide having at least two cysteine
CC residues, and a second gene encoding a positively charged basic
CC antimicrobial peptide. A host microorganism is transformed with a vector
CC containing the fusion gene and then cultured. The expressed antimicrobial
CC peptide is then recovered. The method is used to mass produce
CC antimicrobial peptides in recombinant microorganisms. The inhibitory
CC effect of the expressed antimicrobial peptide upon the growth of the host
CC microorganism is considerably reduced by fusing it to the acidic peptide.
CC Therefore, the use of the fusion gene provides an economic, recombinant
CC alternative of mass producing antimicrobial peptides, which overcomes the
CC disadvantages of low-productivity and poor economy, previously
CC encountered by recombinant and chemical methods. The present sequence
CC represents the DNA encoding an antimicrobial peptide indolicidin. This
CC can be used along with the acidic peptide guamerin gene in the
CC construction of the fusion gene.
XX
SQ Sequence 39 BP; 4 A; 10 C; 16 G; 9 T; 0 other;

alignment_scores:
      Quality: 73.00      Length: 9
      Ratio: 8.111      Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AAV83788

Align seg 1/1 to: AAV83788 from: 1 to: 39

4 LysTrpProTrrTPProTrrPARgArg 12
|||||
13 AATGCGCGTGTGCGCGTGTGCGTGTG 39

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ29389
seq_documentation_block:
ID AAZ29389 standard; DNA; 47 BP.
XX
XX AAZ29389;
AC
XX
DT 29-FEB-2000 (first entry)
XX
DE PCR primer-15 for synthesis of antimicrobial peptide indolicidin gene.
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
OS Synthetic.
XX
XX MO9964611-A1.
PN
XX

```

```

PD 16-DEC-1999.
XX
XX 08-JUN-1999; 99MO-KR00282.
PF
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
XX
PI Kim JH, Kang MH, Lee J, Park SH, Lee JW, Hong SS, Lee H;
XX WPI, 2000-097542/08.
DR
XX
XX New DNA constructs useful for mass production of antimicrobial peptides
PT in microorganism hosts -
PT
XX
PS Example 1; Page 13; 67pp; English.
XX
CC The present sequence is a chemically synthesised PCR primer which was
CC used to synthesise a gene encoding antimicrobial peptide indolicidin.
CC The antimicrobial peptide gene is used in a DNA construct that comprises
CC entire, partial or a derivative of purf gene (glutamine
CC pyrophosphoribosyl pyrophosphate amidotransferase gene). The DNA
CC construct allows mass production of the antimicrobial peptide in
CC microbial hosts without killing the host cells. The antimicrobial
CC peptides are useful commercially in the pharmaceutical and
CC food industries.
XX
SQ Sequence 47 BP; 6 A; 11 C; 19 G; 11 T; 0 other;

alignment_scores:
      Quality: 73.00      Length: 9
      Ratio: 8.111      Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x AAZ29389

Align seg 1/1 to: AAZ29389 from: 1 to: 47

4 LysTrpProTrrTPProTrrPARgArg 12
|||||
17 AATGCGCGTGTGCGCGTGTGCGTGTG 43

seq_name: /SID8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAZ29390
seq_documentation_block:
ID AAZ29390 standard; DNA; 47 BP.
XX
XX AAZ29390;
AC
XX
DT 29-FEB-2000 (first entry)
XX
DE PCR primer-16 for synthesis of antimicrobial peptide indolicidin gene.
XX
XX PCR primer: anti-microbial peptide; indolicidin gene; DNA construct;
KW glutamine pyrophosphoribosyl pyrophosphate amidotransferase gene;
KW purf gene; fusion peptide; mass production; pharmaceutical industry;
KW food industry; ss.
XX
OS Synthetic.
XX
XX MO9964611-A1.
PN
XX
XX 16-DEC-1999.
PD
XX
XX 08-JUN-1999; 99MO-KR00282.
PF
XX
XX 09-JUN-1998; 98KR-0022117.
PR 14-MAY-1999; 99KR-0017920.
XX
XX (SAMY-) SAMYANG GENEX CORP.
PA

```


KM Indolicidin analogue; antimicrobial activity; helminth; bacteria; virus;
 KM treatment; inhibit growth; micro-organism; contact lens solution;
 KM transgenic plant; surgical instrument; yeast; fungi; protozoa; ss.
 XX
 OS Synthetic.
 XX
 PN WO958141-A1.
 XX
 PD 18-NOV-1999.
 XX
 PF 05-MAY-1999; 99WO-US09942.
 XX
 PR 12-MAY-1998; 98US-0076227.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PI Selected ME;
 XX
 DR WPI: 2000-053028/04.
 DR P-SD8; AAY57142.
 XX
 PT New indolicidin analogues, active against bacteria, yeast, fungi,
 PT protozoa and virus, used for, e.g. treating infections -
 XX
 PS Disclosure; Fig 6; 62pp; English.
 XX
 CC This is the nucleotide sequence of an example of a fusion protein which
 CC consists of an indolicidin analogue linked to another peptide.
 CC Peptides AAY57109-157138 and AAY57143-157144 are new indolicidin
 CC analogues, which have a homoserine residue and/or a truncated amino
 CC terminal region. The analogues have the following amino acid sequence:
 CC Xaa1-Xaa2-Xaa3-Xaa4-Xaa5-Xaa6-Pro-Xaa6-Pro-Xaa6-Xaa7-Xaa7-Xaa8
 CC where:
 CC Xaa1 = Ile, Leu, Val, Ala, Gly or absent;
 CC Xaa2 = Ile, Leu, Val, Ala, Gly or absent;
 CC Xaa3 = Pro or absent;
 CC Xaa4 = Trp, Phe or absent;
 CC Xaa5 = Arg, Lys or absent;
 CC Xaa6 = Trp or Phe;
 CC Xaa7 = Arg, Lys or absent;
 CC Xaa8 = homoserine (Hse), Met, Met-Xaa9-Met or absent, and
 CC Xaa9 = at least one amino acid;
 CC provided that if Xaa1 is present, Xaa8 = Hse, Met or Met-Xaa9-Met;
 CC and further provided that: if Xaa2 is absent, Xaa1 is absent; if Xaa3 is
 CC absent, Xaa1 and Xaa2 are absent; if Xaa4 is absent, Xaa1, Xaa2 and Xaa3
 CC are absent; and if Xaa5 is absent, Xaa1, Xaa2, Xaa3 and Xaa4 are absent.
 CC The indolicidin analogues can be used to create a fusion polypeptide
 CC consisting of the analogue linked to a peptide. The indolicidin
 CC analogues have antimicrobial activity against gram positive bacteria,
 CC gram negative bacteria, yeast, fungus, protozoa and viruses (e.g. HIV-1).
 CC They are also active against helminths. The analogues can be used for
 CC reducing or inhibiting growth or survival of a microorganism. They can be
 CC used for treating infections. They can also be included in a liquid such
 CC as water or an aqueous solution, e.g. contact lens solution. The
 CC analogues have potential uses in food products, and in objects such as
 CC the surface of an instrument used to prepare food or to perform surgery.
 CC Transgenic plants or animals useful in the food industry can be produced
 CC by introducing a nucleic acid molecule encoding an indolicidin analogue
 CC into the germ-line cells of such organisms.
 XX
 SQ Sequence 211 BP; 36 A; 50 C; 74 G; 51 T; 0 other;

alignment_scores:
 Quality: 73.00 Length: 9
 Ratio: 8.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
 US-09-444-281-35 x AAZ45123 ..

Align seg 1/1 to: AAZ45123 from: 1 to: 211

4 LysTrpProTrpTrpProTrpArg 12
 |||||
 38 AATGCGCGGTGGTGGCGGTGCT 64

seq_name: /SIDS8/gcgdata/geneseq/geneseq/NA1999.DAT:AAZ20646

seq_documentation_block:
 ID AAZ20646 standard; RNA: 6446 BP.

XX
 AC AAZ20646;
 XX
 DT 26-NOV-1999 (first entry)
 XX
 DE TMV-based virus TMV861 coat protein read-through RNA sequence.
 XX
 KM TMV-based virus; tobacco mosaic virus; protein isolation; green juice;
 KM virus isolation; fusion protein identification; ss.
 XX
 OS Tobacco mosaic virus.
 XX
 PN WO9946288-A2.
 PD 16-SEP-1999.
 XX
 PF 09-MAR-1999; 99WO-US05056.
 XX
 PR 10-MAR-1998; 98US-0037751.
 XX
 PA (BIOS-) BIOSOURCE TECHNOLOGIES INC.
 XX
 PI Garger SJ, Holtz RB, McCulloch MJ, Turpen TH;
 XX WPI: 1999-561660/47.
 DR
 XX
 PT Obtaining protein, viruses and fusion proteins from plants, using
 PT non-denaturing conditions -
 XX
 PS Disclosure; Page 55-58; 58pp; English.
 XX
 CC This sequence represents a tobacco mosaic virus (TMV) based virus
 CC sequence identified using the method of the invention. The method is for
 CC obtaining a soluble protein or peptide of interest from a plant,
 CC comprising homogenising the plant to produce green juice, adjusting the
 CC pH to less than or equal to 5.2, and heating the juice to a minimum of
 CC 45 degrees C. The juice is then centrifuged to produce a supernatant, and
 CC the protein or peptide is purified from the supernatant. The method can
 CC also be used for obtaining viruses and fusion proteins. The method is
 CC especially useful for obtaining IL-1 to IL-10, EPO, G-CSF, GM-CSF,
 CC hp-CSF, M-CSF, Factor VIII, Factor IX, tPA, receptors, receptor
 CC antagonists, antibodies, single-chain antibodies, enzymes,
 CC neuropeptides, insulin, antigens, vaccines, peptide hormones,
 CC calcitonin, and human growth hormone, or an antimicrobial peptide or
 CC protein from protegrins, magainins, cecropins, melittins, indolicidins,
 CC defensins, beta-defensins, cryptidins, clavalins, plant defensins,
 CC nicin and bactericins, all produced by recombinant means. The new method
 CC is more efficient than the prior art for isolating viruses, protein, and
 CC peptides. The method is large-scale, and non-denaturing and
 CC solvent-limited. Prior art methods do not isolate recombinant proteins,
 CC and do not allow fraction 2 proteins to be ultrafiltrated.
 XX
 SQ Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores:
 Quality: 73.00 Length: 9
 Ratio: 8.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
 US-09-444-281-35 x AAZ20646 ..

Align seg 1/1 to: AAZ20646 from: 1 to: 6446

4 LysTrpProTrpTrpParGarg 12
 |||||
 6213 AAGUGCCUUGUGGCCAUGGCCCGA 6239

seq_name: /SIDS8/gcdata/geneseq/geneseqn/NA2001.DAT:AAF82334

seq_documentation_block:

ID AAF82334 standard; RNA; 6446 BP.

AAF82334;

22-JUN-2001 (first entry)

Tobacco mosaic virus-based coat protein read-through virus TMV861.

TMV; tobacco mosaic virus; TMV861; virus isolation;

non-native protein purification; ribulose 1,5-diphosphate carboxylase;

Rubisco; coat protein read-through; ss.

Tobacco mosaic virus.

WO200119969-A1.

22-MAR-2001.

19-MAY-2000; 2000WO-US13680.

16-SEP-1999; 99US-0397090.

(LARG-) LARGE SCALE BIOLOGY CORP.

Garger SJ, Holtz BR, McCulloch MJ, Turpen TH;

WPI; 2001-328016/34.

Minimizing presence of ribulose 1,5-diphosphate carboxylase to obtain

plant product for isolating bioactive species involves cutting plant

material from plant in cutting period when quantity of Rubisco is at

minimum

Disclosure; Page 72-74; 81pp; English.

The present sequence is a tobacco mosaic virus (TMV)-based virus

which was used to infect field-grown tobacco. The virus was then

isolated from the tobacco plants by a novel process for isolating and

purifying viruses, soluble proteins and peptides from plant sources. In

order to isolate the bioactive species from the undesirable

photosynthetic protein ribulose 1,5-diphosphate carboxylase (Rubisco),

the plant material is cut in a period of the light/dark cycle when the

quantity of Rubisco in the plant is at a minimum. The method is useful

for obtaining a virus of interest. It is also useful for obtaining

soluble recombinant or non-native proteins, such as active mammalian

proteins, enzymes, vaccines, antibodies and peptides, from transgenic

plants.

Sequence 6446 BP; 1873 A; 1234 C; 1563 G; 1776 U; 0 other;

alignment_scores: Quality: 73.00 Length: 9

Ratio: 8.111 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x AAF82334 ..

Align seg 1/1 to: AAF82334 from: 1 to: 6446

4 LysTrpProTrpTrpParGarg 12

|||||
 6213 AAGUGCCUUGUGGCCAUGGCCCGA 6239

seq_name: /SIDS8/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28519

seq_documentation_block:

ID AAA28519 standard; DNA; 207 BP.

AAA28519;

29-AUG-2000 (first entry)

PCRIL DNA coding sequence.

Magalain; antimicrobial; transgenic plant; protease degradation; Rev4;

indolicidin; protein production; reverse peptide; ss.

Synthetic.

WO200026344-A1.

11-MAY-2000.

29-OCT-1999; 99WO-US25561.

30-OCT-1998; 98US-0106373.

02-NOV-1998; 98US-0106537.

(INTE-) INTERLINK BIOTECHNOLOGIES LLC.

(KENT) UNIV KENTUCKY RES FOUND.

Everett NP, Li Q, Lawrence C, Davies MH;

WPI; 2000-365597/31.

P-PSDB; AAY92840.

Polypeptides for reducing proteolytic degradation of proteins

administered to, or produced by a plant comprise indolicidin or its

functional equivalents

Example 17; Page 35; 50pp; English.

Indolicidin is a potent antimicrobial tripeptide, originally

purified from cytoplasmic granules of bovine neutrophils. Reverse

peptide, Rev4 of indolicidin (see AAY92794) was found to have increased

stability against plant protease degradation. Expression of antimicrobial

peptides in transgenic plants suffers a major limitation in that the

foreign peptides are susceptible to rapid degradation by proteases. The

invention concerns reducing the extent of protease degradation of a

protein applied to, or produced by a plant by administering indolicidin,

Rev4 or a functional equivalent to the plant. Transgenic plants

expressing indolicidin and Rev4 are useful for production of the

antimicrobial peptides. Compositions containing indolicidin and Rev4 are

also useful for production of agronomically important proteins in

plants.

Sequence 207 BP; 49 A; 50 C; 36 G; 72 T; 0 other;

alignment_scores: Quality: 66.00 Length: 10

Ratio: 6.600 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 60.000

alignment_block:

US-09-444-281-35 x AAA28519 ..

Align seg 1/1 to: AAA28519 from: 1 to: 207

2 LeuLysLysTrpProTrpTrpParG 11

|||||
 163 ATTAGAGATGCGCTTGTGGCTTGAAA 192

APPLICANT: Kim, Jeong Hyun
APPLICANT: Hong, Seung-Suh
APPLICANT: Lee, Hyun-Soo
APPLICANT: Samsung Genex Corporation
TITLE OF INVENTION: METHOD FOR MASS PRODUCTION OF
TITLE OF INVENTION: ANTIMICROBIAL PEPTIDE
FILE REFERENCE: 6181/0F135
CURRENT APPLICATION NUMBER: US/09/230,180
CURRENT FILING DATE: 1999-03-10
PRIOR APPLICATION NUMBER: PCT/KR98/00132
PRIOR FILING DATE: 1998-05-28
PRIOR APPLICATION NUMBER: KR 13372/1998
PRIOR FILING DATE: 1998-04-09
PRIOR APPLICATION NUMBER: KR 21312/1997
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 36
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 29
LENGTH: 39
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: DNA sequence deduced from Indolicidin peptide
US-09-230-180-29

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x US-09-230-180-29
Align seg 1/1 to: US-09-230-180-29 from: 1 to: 39

4 LysTrpProTirPTrProtParGarg 12
|||||
13 AATGCGCGTGGCGCGCGTGGT 39

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-259-741-5

seq_documentation_block:

Sequence 5, Application US/09259741
Patent No. 6033895
GENERAL INFORMATION:
APPLICANT: GARGER, STEPHEN
APPLICANT: HOLTZ, R. BARRY
APPLICANT: MCCULLOCH, MICHAEL
APPLICANT: TURPEN, THOMAS
TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES FROM PLANT
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howrey & Simon
STREET: 1299 Pennsylvania Avenue N.W.
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/259,741
FILING DATE: February 25, 1999
CLASSIFICATION:
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/037,751
FILING DATE: March 10, 1998
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801.0140.US01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8100
TELEFAX: 650-463-8400
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: Genomic RNA
US-09-259-741-5

alignment_scores:
Quality: 73.00 Length: 9
Ratio: 8.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x US-09-259-741-5
Align seg 1/1 to: US-09-259-741-5 from: 1 to: 6446

4 LysTrpProTirPTrProtParGarg 12
|||||
6213 AAGUGCCUUGGCGCAGCGCCGA 6239

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-09-037-751-5

seq_documentation_block:

Sequence 5, Application US/09037751
Patent No. 6037456
GENERAL INFORMATION:
APPLICANT: GARGER, STEPHEN
APPLICANT: HOLTZ, R. BARRY
APPLICANT: MCCULLOCH, MICHAEL
APPLICANT: TURPEN, THOMAS
TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
TITLE OF INVENTION: PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Howrey & Simon
STREET: 1299 Pennsylvania Avenue N.W.
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20004
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/037,751
FILING DATE: 10-MAR-1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801.0140.999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8109

```

; TELEFAX: 650-463-8400
; TELEX:
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6446 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: unknown
; MOLECULE TYPE: Genomic RNA
; US-09-037-751-5

alignment_scores:
    Quality: 73.00      Length: 9
    Ratio: 8.111      Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-037-751-5 ..

Align seg 1/1 to: US-09-037-751-5 from: 1 to: 6446

4 LysTrpProTTrpTTrpTTrpArgArg 12
|||||
6213 AAGUGCCUGGUGGCGCAUGGCGCGCA 6239

seq_name: /cgn2_6/prodata/2/lna/6B_COMB.seq:US-09-466-422-5

seq_documentation_block:
; Sequence 5, Application US/09466422
; Patent No. 6303779
; GENERAL INFORMATION:
; APPLICANT: GARGER, STEPHEN
; HOLTZ, R. BARRY
; MCCULLOCH, MICHAEL
; TURPEN, THOMAS
; TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
; PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES
; FROM PLANT SOURCES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Howrey & Simon
; STREET: 1299 Pennsylvania Avenue N.W.
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20004
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTED for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/466,422
; FILING DATE: 17-Dec-1999
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/037,751
; FILING DATE: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Halluin, Albert P
; REGISTRATION NUMBER: 25,277
; REFERENCE/DOCKET NUMBER: 00801.0140.999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-463-8109
; TELEFAX: 650-463-8400
; TELEX: <Unknown>
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6446 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: unknown
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; MOLECULE TYPE: Genomic RNA
; SEQUENCE DESCRIPTION: SEQ ID NO: 5:
; US-09-466-422-5

alignment_scores:
    Quality: 73.00      Length: 9
    Ratio: 8.111      Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 100.000

alignment_block:
US-09-444-281-35 x US-09-466-422-5 ..

Align seg 1/1 to: US-09-466-422-5 from: 1 to: 6446

4 LysTrpProTTrpTTrpTTrpArgArg 12
|||||
6213 AAGUGCCUGGUGGCGCAUGGCGCGCA 6239

seq_name: /cgn2_6/prodata/2/lna/5A_COMB.seq:US-08-159-784-1

seq_documentation_block:
; Sequence 1, Application US/08159784
; Patent No. 5643783
; GENERAL INFORMATION:
; APPLICANT: Bjorn R. Olsen
; TITLE OF INVENTION: NOVEL COLLAGEN AND USES THEREOF
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; COMPUTER: IBM PS/2 Model 502 or 558x
; OPERATING SYSTEM: MS-DOS (Version 5.0)
; SOFTWARE: WordPerfect (Version 5.1)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/159,784
; FILING DATE: December 1, 1993
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: John F. Freeman
; REGISTRATION NUMBER: 29,066
; REFERENCE/DOCKET NUMBER: 00246/170001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 542-5070
; TELEFAX: (617) 542-8906
; TELEX: 200154
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 4031
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-159-784-1

alignment_scores:
    Quality: 60.00      Length: 9
    Ratio: 7.500      Gaps: 0
    Percent Similarity: 88.889    Percent Identity: 77.778

alignment_block:
US-09-444-281-35 x US-08-159-784-1/rev ..

Align seg 1/1 to reverse of: US-08-159-784-1 from: 1 to: 4031
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3 LysLysTrpProTrpProTrp 11
|||||
1671 AATCCTGCTGCTGCTGCTG 1645

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-481-337A-1

seq_documentation_block:

Sequence 1, Application US/08481337A
Patent No. 5863738
GENERAL INFORMATION:
APPLICANT: TEN DUKE, Peter
APPLICANT: HELDIN, Carl-Henrik
APPLICANT: MIYAZONO, Konei
APPLICANT: SAMPATH, Kuber T.
TITLE OF INVENTION: Morphogenic Protein-Specific Cell
TITLE OF INVENTION: Surface Receptors and Uses Therefor
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: Testa, Hurwitz & Thibault
STREET: 125 High St.
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02110
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/481,337A
FILING DATE: 02-JUN-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: MEYERS, Thomas C.
REGISTRATION NUMBER: 36,989
REFERENCE/DOCKET NUMBER: CRP-097CP2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1509 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1509
OTHER INFORMATION: /product= "Human ALK1"
US-08-481-337A-1

alignment_scores:

Quality:	58.00	Length:	6
Ratio:	9.667	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	100.000

alignment_block:

US-09-444-281-35 x US-08-481-337A-1

Align seg 1/1 to: US-08-481-337A-1 from: 1 to: 1509

5 TrpProTrpProTrp 10

|||||
389 TGCCCTGCTGCTGCTG 406

seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-696-268B-1

seq_documentation_block:

Sequence 1, Application US/08696268B

Patent No. 5968752

GENERAL INFORMATION:

APPLICANT: ICHIO, Hidenori

APPLICANT: NISHIOH, Hidenori

APPLICANT: SAMPATH, Kuber T.

TITLE OF INVENTION: NOVEL SIGNALING RECEPTOR FOR

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Testa, Hurwitz & Thibault

STREET: 125 High St.

CITY: Boston

STATE: MA

COUNTRY: USA

ZIP: 02110

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/696,268B

FILING DATE:

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: MEYERS, Thomas C.

REGISTRATION NUMBER: 36,989

REFERENCE/DOCKET NUMBER: CRP-117

TELECOMMUNICATION INFORMATION:

TELEPHONE: (617) 248-7000

TELEFAX: (617) 248-7100

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1509 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

FEATURE:

NAME/KEY: CDS

LOCATION: 1..1509

OTHER INFORMATION: /product= "Human ALK-1"

US-08-696-268B-1

alignment_scores:

Quality:	58.00	Length:	6
Ratio:	9.667	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	100.000

alignment_block:

US-09-444-281-35 x US-08-696-268B-1

Align seg 1/1 to: US-08-696-268B-1 from: 1 to: 1509

5 TrpProTrpProTrp 10

|||||
389 TGCCCTGCTGCTGCTG 406

seq_name: /cgn2_6/ptodata/2/ina/PCTUS_COMB.seq:PCT-US95-05467-1

seq_documentation_block:

Sequence 1, Application PC/TUS9505467

GENERAL INFORMATION:

APPLICANT:

APPLICANT:

TITLE OF INVENTION: MORPHOGENIC PROTEIN-SPECIFIC CELL

TITLE OF INVENTION: SURFACE RECEPTORS AND USES THEREFOR

NUMBER OF SEQUENCES: 15

CORRESPONDENCE ADDRESS:

ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &

STREET: 53 STATE STREET

CITY: BOSTON
STATE: MA
COUNTRY: USA
ZIP: 02109
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/05467
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: PITCHER, EDMUND R.
REGISTRATION NUMBER: 27,829
REFERENCE/DOCKET NUMBER: CRP-097PC
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ. ID NO.: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1509 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1509
OTHER INFORMATION: /product= "Human ALK1"
PCT-US95-05467-1

alignment_scores:
Quality: 58.00 Length: 6
Ratio: 9.667 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x PCT-US95-05467-1 ..

Align seg 1/1 to: PCT-US95-05467-1 from: 1 to: 1509

5 TrpProTrrTrrProTrr 10
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389 TGCCCTGTGTGCTCG 406

seq_name: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:PCT-US94-11328A-3

seq_documentation_block:

Sequence 3, Application PC/TUS9411328A
GENERAL INFORMATION:
APPLICANT: HE, ET AL.
TITLE OF INVENTION: TAR-1 and TAR-3
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: CARELLA, HYRNE, BAIN, GILFILLAN,
ADDRESS: CECCHI, STEWART & OLSTEIN
STREET: 6 BECKER FARM ROAD
CITY: ROSELAND
STATE: NEW JERSEY
COUNTRY: USA
ZIP: 07068
COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5 INCH DISKETTE
COMPUTER: IBM PS/2
OPERATING SYSTEM: MS-DOS
SOFTWARE: WORD PERFECT 5.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US94/11328A
FILING DATE: Submitted herewith
CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: FERRARO, GREGORY D.
REGISTRATION NUMBER: 36,134
REFERENCE/DOCKET NUMBER: 325800-132
TELECOMMUNICATION INFORMATION:
TELEPHONE: 201-994-1700
TELEFAX: 201-994-1744
INFORMATION FOR SEQ. ID NO.: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1596 BASE PAIRS
TYPE: NUCLEIC ACID
STRANDEDNESS: SINGLE
TOPOLOGY: LINEAR
MOLECULE TYPE: cDNA
PCT-US94-11328A-3

alignment_scores:
Quality: 58.00 Length: 6
Ratio: 9.667 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x PCT-US94-11328A-3 ..

Align seg 1/1 to: PCT-US94-11328A-3 from: 1 to: 1596

5 TrpProTrrTrrProTrr 10
|||||
470 TGCCCTGTGTGCTCG 487

seq_name: /cgn2_6/prodata/2/ina/GB_COMB.seq:US-09-382-256-1

seq_documentation_block:

Sequence 1, Application US/09382256A
Patent No. 6207814
GENERAL INFORMATION:
APPLICANT: MIYAZONO, Kohel
TEN DIJKE, Peter
FRANZEN, Petra
YAMASHITA, Hideohshi
HELDIN, Carl-Henrik
TITLE OF INVENTION: ACTIVIN RECEPTOR LIKE KINASES, PROTEINS
HAVING SERINE THREONINE KINASE DOMAINS,
AND THEIR USE
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: Fulbright & Jaworski L.L.P.
STREET: 666 Fifth Avenue
CITY: New York City
STATE: New York
COUNTRY: USA
ZIP: 10103
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.25 inch, 1.44mb
COMPUTER: IBM PS/2
OPERATING SYSTEM: PC-DOS
SOFTWARE: Wordperfect
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/382,256A
FILING DATE: 24-Aug-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/GB93/02367
FILING DATE: No. 6207814 member 17, 1993
APPLICATION NUMBER: GB 9224057.1
FILING DATE: No. 6207814 member 17, 1992
APPLICATION NUMBER: GB 9304677.9
FILING DATE: March 8, 1993
APPLICATION NUMBER: GB 9304680.3

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; FILING DATE: March 8, 1993
; APPLICATION NUMBER: 9311047.6
; FILING DATE: May 28, 1993
; APPLICATION NUMBER: 9313763.6
; FILING DATE: July 2, 1993
; APPLICATION NUMBER: 9316099.2
; FILING DATE: August 3, 1993
; APPLICATION NUMBER: 321344.5
; FILING DATE: October 15, 1993
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6207814man D. Hanson
; REGISTRATION NUMBER: 30,946
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 318-3000
; TELEFAX: (212) 752-5958
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1984 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: Internal
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 283..1791
; US-09-382-256-1
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:

alignment_scores:
    Quality: 58.00      Length: 6
    Ratio: 9.667      Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 100.000

alignment_block:
    US-09-444-281-35 x US-09-382-256-1
Align seg 1/1 to: US-09-382-256-1 from: 1 to: 1984
    5 TrpProTrrpProTrrp 10
    ||||||||||||||||
    671 TGGCCCTGTGGCTGG 688

seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-395-115-1
seq_documentation_block:
; Sequence 1, Application US/09395115
; Patent No. 6271365
; GENERAL INFORMATION:
; APPLICANT: Miyazono, Kohel; Dijke, Peter Ten;
; TITLE OF INVENTION: Activin Receptor-Like Kinase, Proteins
; TITLE OF INVENTION: Having Serine Threonine Kinase Domains And Their Use
; NUMBER OF SEQUENCES: 29
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Felfe & Lynch
; STREET: 805 Third Avenue
; CITY: New York City
; STATE: New York
; ZIP: 10022
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 360 kb storage
; COMPUTER: IBM
; OPERATING SYSTEM: PC-DOS
; SOFTWARE: Wordperfect
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/395,115
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; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/436,265
; FILING DATE: 30-October-1995
; APPLICATION NUMBER: PCT/GB93/02367
; FILING DATE: 17-No. 6271365ember-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9224057.1
; FILING DATE: 17-No. 6271365ember-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9304677.9
; FILING DATE: 8-March-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9304680.3
; FILING DATE: 8-March-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9311047.6
; FILING DATE: 28-May-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9313763.6
; FILING DATE: 2-July-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9136099.2
; FILING DATE: 3-August-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 9321344.5
; FILING DATE: 15-October-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Kohel, Vineet
; REGISTRATION NUMBER: 37,003
; REFERENCE/DOCKET NUMBER: LUD 5298
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 688-9200
; TELEFAX: (212) 838-3884
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1984 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: unknown
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: Internal
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 283..1791
; US-09-395-115-1

alignment_scores:
    Quality: 58.00      Length: 6
    Ratio: 9.667      Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 100.000

alignment_block:
    US-09-444-281-35 x US-09-395-115-1
Align seg 1/1 to: US-09-395-115-1 from: 1 to: 1984
    5 TrpProTrrpProTrrp 10
    ||||||||||||||||
    671 TGGCCCTGTGGCTGG 688

seq_name: /cgn2_6/ptodata/2/ina/6A_COMB.seq:US-08-749-816-1
seq_documentation_block:
; Sequence 1, Application US/08749816
; Patent No. 6013470
; GENERAL INFORMATION:
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Mon Jan 7 10:42:11 2002

us-09-444-281-35.rni

Page 8

Ratio: 6.389 Gaps: 1
Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:
US-09-444-281-35 x US-08-149-097D-24 ...

Align seg 1/1 to: US-08-149-097D-24 from: 1 to: 7032

2 LeuLysLysTrpProTrp...TrpProTrpArg 11
|||||: ||||| ||||| ||||| |||||
2606 TTGAGGGCCTGGCCCTGGCCCTGGCCCTGGAGA 2638

seq_name: /cgn2.6/ptodata/2/ina/6A.COMB.seq:US-08-949-386-24

seq_documentation_block:
; Sequence 24, Application US/08949386
; Patent No. 6090623
; GENERAL INFORMATION:
; APPLICANT: Harpold, Michael
; APPLICANT: Ellis, Steven
; APPLICANT: Williams, Mark
; APPLICANT: McCue, Ann
; APPLICANT: Gillespie, Allison
; TITLE OF INVENTION: HUMAN CALCIUM CHANNEL COMPOSITIONS AND
; TITLE OF INVENTION: METHODS
; NUMBER OF SEQUENCES: 38
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brown, Martin, Haller & McClain
; STREET: 1660 Union Street
; CITY: San Diego
; STATE: California
; COUNTRY: US
; ZIP: 92101
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/949,386
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/290,012
; FILING DATE: 11-AUG-1994
; APPLICATION NUMBER: 08/149,097
; FILING DATE: 5-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/105,536
; FILING DATE: 11-AUG-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Seidman, Stephanie L.
; REGISTRATION NUMBER: 33,779
; REFERENCE/DOCKET NUMBER: 519808
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 238-0999
; TELEFAX: (619) 238-0062
; INFORMATION FOR SEQ ID NO: 24:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7032 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 166..6921
; OTHER INFORMATION: /standard_name="Alpha-1E-1"
US-08-949-386-24

alignment_scores:
Quality: 57.50 Length: 11

Ratio: 6.389 Gaps: 1
Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:
US-09-444-281-35 x US-08-949-386-24 ..

Align seg 1/1 to: US-08-949-386-24 from: 1 to: 7032

2 LeuLysLysTrpProTrp...TrpProTrpArg 11
|||||: ||||| ||||| ||||| |||||
2606 TTGAGGGCCTGGCCCTGGCCCTGGCCCTGGAGA 2638


```

ID  AAV90541 standard; cDNA; 375 BP.
XX
AC  AAV90541;
XX
DE  15-FEB-1999 (first entry)
XX
DE  EST clone BK517.
XX
KM  Human; secreted protein; expressed sequence tag; EST; haematopoiesis;
KM  tissue growth; activin; inhibin; chemotaxis; chemokinesis; haemostatic;
KM  receptor; ligand; thrombolytic; anti-inflammatory; cadherin; anti-tumour;
KM  gene therapy; ss.
XX
OS  Homo sapiens.
XX
PN  M09845436-A2.
XX
PD  15-OCT-1998.
XX
PF  10-APR-1998; 98MO-US06955.
XX
PR  10-APR-1997; 97US-0838821.
XX
PA  (GEM ) GENETICS INST INC.
XX
PI  Agostino MJ, Jacobs K, Lavallie ER, McCoy JM, Merberg D;
PI  Racie LA, Spaulding V, Treacy M;
XX
DR  MPI; 1999-070077/06.
XX
PT  New polynucleotides encoding human secreted proteins - derived from
PT  e.g. human blood, kidney, foetal lung, placenta, testes, brain,
PT  ovary, pituitary, retina and colon cDNA libraries.
XX
PS  Claim 1; Page 574; 618p; English.
XX
CC  The present sequence represents a human expressed sequence tag (EST).
CC  The polynucleotide, which is a secreted EST, and the encoded protein
CC  are predicted to have useful biological activities which would make
CC  them suitable for treating, preventing or ameliorating medical
CC  conditions in humans and animals, although no supporting data is
CC  given. Suggested activities include nutritional activity, immune
CC  stimulating or suppressing activity, haematopoiesis regulating
CC  activity, tissue growth activity, activin/inhibin activity,
CC  chemotactic/chemokinetic activity, haemostatic and thrombolytic
CC  activity, receptor/ligand activity, anti-inflammatory activity,
CC  cadherin/tumour invasion suppressor activity, tumour inhibition
CC  activity. The polynucleotide may also be useful for gene therapy.
XX
SQ  Sequence 375 BP; 56 A; 125 C; 113 G; 81 T; 0 other;

alignment_scores:
    Quality: 41.00      Length: 9
    Ratio: 4.556       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAV90541/rev ..

Align seg 1/1 to reverse of: AAV90541 from: 1 to: 375

1 HisGluAlaGluProGluAlaGluPro 9
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140 CACGAAGCAGAGCCTGAGGGAGCCG 114

seq_name: /SID88/gcdata/geneseq/geneseqn/NA2001.DAT:AAH10202
seq_documentation_block:
ID  AAH10202 standard; cDNA; 557 BP.
XX
AC  AAH10202;
XX

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DT  26-JUN-2001 (first entry)
XX
DE  Human cDNA clone (3'-primer) SEQ ID NO:7037.
XX
DE  Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS  Homo sapiens.
XX
PN  EP1074617-A2.
XX
PD  07-FEB-2001.
XX
PF  28-JUL-2000; 2000EP-0116126.
XX
PR  29-JUL-1999; 99JP-0248036.
PR  27-AUG-1999; 99JP-0300253.
PR  11-JAN-2000; 2000JP-0118776.
PR  02-MAY-2000; 2000JP-0183767.
PR  09-JUN-2000; 2000JP-0241899.
XX
PA  (HELI-) HELIX RES INST.
XX
PI  Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
PI  Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX
DR  MPI; 2001-318749/34.
XX
PT  Primer sets for synthesizing polynucleotides, particularly the 5602
PT  full-length cDNAs defined in the specification, and for the detection
PT  and/or diagnosis of the abnormality of the proteins encoded by the
PT  full-length cDNAs -
XX
PS  Claim 3; SEQ ID 7037; 2537pp + CD ROM; English.
XX
CC  The present invention describes primer sets for synthesizing 5602
CC  full-length cDNAs defined in the specification. Where a primer set
CC  comprises: (a) an oligo-dr primer and an oligonucleotide complementary
CC  to the complementary strand of a polynucleotide which comprises one of
CC  the 5602 nucleotide sequences defined in the specification, where the
CC  oligonucleotide comprises at least 15 nucleotides; or (b) a combination
CC  of an oligonucleotide comprising a sequence complementary to the
CC  complementary strand of a polynucleotide which comprises a 5'-end
CC  sequence and an oligonucleotide comprising a sequence complementary to a
CC  polynucleotide which comprises a 3'-end sequence, where the
CC  oligonucleotide comprises at least 15 nucleotides and the combination of
CC  the 5'-end sequence/3'-end sequence is selected from those defined in
CC  the specification. The primer sets can be used in antisense therapy and
CC  in gene therapy. The primers are useful for synthesizing polynucleotides,
CC  particularly full-length cDNAs. The primers are also useful for the
CC  detection and/or diagnosis of the abnormality of the proteins encoded by
CC  the full-length cDNAs. The primers allow obtaining of the full-length
CC  cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC  AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
CC  AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC  represent oligonucleotides, all of which are used in the exemplification
CC  of the present invention.
XX
SQ  Sequence 557 BP; 99 A; 165 C; 183 G; 103 T; 7 other;

alignment_scores:
    Quality: 41.00      Length: 9
    Ratio: 4.556       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAH10202 ..

Align seg 1/1 to: AAH10202 from: 1 to: 557

1 HisGluAlaGluProGluAlaGluPro 9
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253 CACGAAGCAGAGCCTGAGGGAGCCG 279

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seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA1991.DAT:AAQ12864
seq_documentation_block:
ID   AAQ12864 standard; DNA: 849 BP.
XX
XX   AAQ12864;
AC
XX   11-OCT-1991 (first entry)
DT
XX   Human Cytotoxic Cell Protease-1 coding sequence.
DE
XX   hccp1 inhibitor; cytotoxic T-lymphocytes; ss.
KM
XX   Homo sapiens.
OS
XX   WO9110685-A.
PN
XX   25-JUL-1991.
PD
XX   17-JAN-1991; 91WO-US00340.
PF
XX   19-JAN-1990; 90US-0467880.
PR
XX   (SERA-) SERAGEN INC.
PA
XX   Bleackley RC, Lobe CG, Paetkau VH, James MN, Murphy M;
PI   WPI; 1991-237989/32.
DR
XX   DNA vectors; and inhibitors of cytotoxic cell protease - for
PT   treatment of auto-immune diseases e.g. pernicious anaemia,
PT   rheumatoid arthritis, allo-graft rejection etc.
PS
XX   Disclosure; Fig 6; 62pp; English.
XX
XX   The hccp1 coding sequence was isolated from cytotoxic T-cell
CC   lymphocytes. Vectors comprising the hccp1 coding sequence are
CC   claimed. Clone hcc1 was isolated and found to be the human analogue
CC   of murine C11.
CC   See AAQ12862-6 and AAR1354-R13262.
CC
XX   Sequence 849 BP; 232 A; 230 C; 222 G; 165 T; 0 other;
SQ
alignment_scores:
Quality: 41.00 Length: 11
Ratio: 4.100 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636
alignment_block:
US-09-444-281-27 x AAQ12864 ..
Align seg 1/1 to: AAQ12864 from: 1 to: 849
1 HisGLUALAGLupProGLUALAGLupProIleMet 11
|||||
68 CATGAGGCCGAGCCCACTCCGCCCTACATG 100
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2000.DAT:AAA28524
seq_documentation_block:
ID   AAA28524 standard; cDNA; 1601 BP.
XX
XX   AAA28524;
AC
XX   29-AUG-2000 (first entry)
DT
XX   Human opioid growth factor receptor cDNA spliced version 1.
DE
XX   OGFfr: opioid growth factor receptor; growth inhibitor; proliferative;
KM   cytostatic; antiviral; gene therapy; antagonist; chromosome 20q13.3; ss.
XX

```

```

OS   Homo sapiens.
XX
XX   Key
FH   5'UTR
FT
XX   CDS
FT
XX   3'UTR
FT
XX   WO200026340-A2.
PN
XX   11-MAY-2000.
PD
XX   02-NOV-1999; 99WO-US25802.
PE
XX   03-NOV-1998; 98US-0106879.
PR
XX   (PENN-) PENN STATE RES FOUND.
PA
XX   Zagon IS, McLaughlin PJ, Verderame MF;
PI   WPI; 2000-365594/31.
DR
XX   P-PSDB; AAY92810.
DR
XX
XX   New cDNA encoding rat and human opioid growth factor receptors which
PT   modulate cell growth, useful for treating cancer
PS   Claim 1; Page 82-83; 91pp; English.
XX
XX   Primers generated from rat opioid growth factor receptor (OGFr) cDNA were
CC   used to clone a fragment of the human OGFr cDNA. The complete sequence of
CC   human OGFr was assembled with a combination of 3' and 5' RACE. 5' RACE
CC   consistently yielded a single species of cDNA, while the 3' RACE revealed
CC   extensive alternative splicing. The alternate splice forms were missing
CC   the imperfect repeats or differed in the number of imperfect repeats. The
CC   human OGFr gene chromosomal location was determined by FISH as 20q13.3.
CC   OGFr proteins, nucleic acid molecules, antibodies, transformed cells and
CC   expression vector are useful for detecting expression or levels of an
CC   OGFr in a tissue. OGFr nucleic acids can be used to inhibit growth of
CC   cells in vitro. The antisense sequences and antibodies can be used to
CC   promote growth of cells in vitro. Cell growth can be promoted by
CC   interfering with the OGFr ligand-receptor system, especially where a
CC   subject suffers from a tissue wound. Treating cancer comprises enhancing
CC   the function of the OGFr ligand-receptor system in cancerous cells of a
CC   patient or administering the OGFr nucleic acid to the patient.
CC
XX   Sequence 1601 BP; 322 A; 485 C; 558 G; 236 T; 0 other;
SQ
alignment_scores:
Quality: 41.00 Length: 9
Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778
alignment_block:
US-09-444-281-27 x AAA28524/rev ..
Align seg 1/1 to reverse of: AAA28524 from: 1 to: 1601
1 HisGLUALAGLupProGLUALAGLupPro 9
|||||
1418 CACGAAGCAGAGCTGAGGAGGCCG 1392
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seq_documentation_block:
ID   AAA28526 standard; cDNA; 2289 BP.
XX
XX   AAA28526;
AC
XX   29-AUG-2000 (first entry)
DT

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XX DE Human opioid growth factor receptor cDNA spliced version 7.
XX XX
XX OGFR: opioid growth factor receptor; growth inhibitor; proliferative;
XX KM cytostatic; vulnerrary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX OS Homo sapiens.
XX FH Key
XX FT 5'UTR 1..33 Location/Qualifiers
XX FT CDS /tag= a
XX FT /tag= b
XX FT /product= opioid_growth_factor_receptor
XX FT 2008..2289
XX FT /tag= c
XX PN WO200026340-A2.
XX PD 11-MAY-2000.
XX PF 02-NOV-1999; 99WO-US25802.
XX PR 03-NOV-1998; 98US-0106879.
XX PR (PENN-) PENN STATE RES FOUND.
XX PI Zagon IS, McLaughlin PJ, Verderame MF;
XX DR MPI: 2000-365594/31.
XX DR P-PSDB: AAY92812.
XX PT New cDNA encoding rat*and human opioid growth factor receptors which
XX PT modulate cell growth, useful for treating cancer
XX PS Claim 1; Page 87-89; 91pp; English.
XX PS
XX CC Primers generated from rat opioid growth factor receptor (OGFR) cDNA were
XX CC used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX CC human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX CC consistently yielded a single species of cDNA, while the 3' RACE revealed
XX CC extensive alternative splicing. The alternate splice forms were missing
XX CC the imperfect repeats or differed in the number of imperfect repeats. The
XX CC human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX CC OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
XX CC expression vector are useful for detecting expression or levels of an
XX CC OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX CC cells in vitro. The antisense sequences and antibodies can be used to
XX CC promote growth of cells in vitro. Cell growth can be promoted by
XX CC interfering with the OGFR ligand-receptor system, especially where a
XX CC subject suffers from a tissue wound. Treating cancer comprises enhancing
XX CC the function of the OGFR ligand-receptor system in cancerous cells of a
XX CC patient or administering the OGFR nucleic acid to the patient.
XX SQ Sequence 2289 BP; 470 A; 714 C; 809 G; 296 T; 0 other;

alignment_scores:
    Quality: 41.00 Length: 9
    Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAA28526/rev ..
Align seg 1/1 to reverse of: AAA28526 from: 1 to: 2289
1 HHSGLUAlaGluProGluAlaGluPro 9
|||||
2105 CACGAAGCAGAGCCTGAGGAGCCCG 2079
seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28522
```

```
seq_documentation_block:
ID AAA28522 standard; cDNA: 2290 BP.
XX AC AAA28522;
XX AC 29-AUG-2000 (first entry)
XX DE Human opioid growth factor receptor cDNA of spliced form A.
XX KM OGFR: opioid growth factor receptor; growth inhibitor; proliferative;
XX KM cytostatic; vulnerrary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX OS Homo sapiens.
XX FH Key
XX FT CDS 34..2007 Location/Qualifiers
XX FT /tag= a
XX PN WO200026340-A2.
XX PD 11-MAY-2000.
XX PF 02-NOV-1999; 99WO-US25802.
XX PR 03-NOV-1998; 98US-0106879.
XX PR (PENN-) PENN STATE RES FOUND.
XX PI Zagon IS, McLaughlin PJ, Verderame MF;
XX DR MPI: 2000-365594/31.
XX DR New cDNA encoding rat and human opioid growth factor receptors which
XX DR modulate cell growth, useful for treating cancer
XX PT Claim 1; Page 77-78; 91pp; English.
XX PS
XX CC Primers generated from rat opioid growth factor receptor (OGFR) cDNA were
XX CC used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX CC human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX CC consistently yielded a single species of cDNA, while the 3' RACE revealed
XX CC extensive alternative splicing. The alternate splice forms were missing
XX CC the imperfect repeats or differed in the number of imperfect repeats. The
XX CC human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX CC OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
XX CC expression vector are useful for detecting expression or levels of an
XX CC OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX CC cells in vitro. The antisense sequences and antibodies can be used to
XX CC promote growth of cells in vitro. Cell growth can be promoted by
XX CC interfering with the OGFR ligand-receptor system, especially where a
XX CC subject suffers from a tissue wound. Treating cancer comprises enhancing
XX CC the function of the OGFR ligand-receptor system in cancerous cells of a
XX CC patient or administering the OGFR nucleic acid to the patient.
XX SQ Sequence 2290 BP; 470 A; 713 C; 807 G; 297 T; 3 other;

alignment_scores:
    Quality: 41.00 Length: 9
    Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x AAA28522/rev ..
Align seg 1/1 to reverse of: AAA28522 from: 1 to: 2290
1 HHSGLUAlaGluProGluAlaGluPro 9
|||||
2106 CACGAAGCAGAGCCTGAGGAGCCCG 2080
seq_name: /SID58/gcdata/geneseq/geneseqn/NA2000.DAT:AAA28525
seq_documentation_block:
```



```

ID AAA28525 standard; cDNA; 2348 BP.
XX
XX AAA28525;
AC
XX 29-AUG-2000 (first entry)
XX
XX Human opioioid growth factor receptor cDNA spliced version 4.
XX
XX OGFR; opioioid growth factor receptor; growth inhibitor; proliferative;
XX cytostatic; vulnerary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX 5'UTR 1..33
XX FT /*tag= a
XX FT 34..2067
XX CDS /*tag= b
XX FT /product= Opioid_growth_factor_receptor
XX FT 2068..2348
XX FT /*tag= c
XX
XX 3'UTR
XX
XX WO200026340-A2.
XX
XX 11-MAY-2000.
XX
XX 02-NOV-1999; 99WO-US25802.
XX
XX 03-NOV-1998; 98US-0106879.
XX
XX (PENN-) PENN STATE RES FOUND.
XX
XX Zagon IS, McLaughlin PJ, Verderame MF;
XX WPI; 2000-365594/31.
XX DR P-PSDB; AAY92811.
XX
XX New cDNA encoding rat and human opioioid growth factor receptors which
XX modulate cell growth, useful for treating cancer
XX
XX Claim 1; Page 84-86; 91pp; English.
XX
XX Primers generated from rat opioioid growth factor receptor (OGFR) cDNA were
XX used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX consistently yielded a single species of cDNA, while the 3' RACE revealed
XX extensive alternative splicing. The alternate splice forms were missing
XX the imperfect repeats, or differed in the number of imperfect repeats. The
XX human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX OGFR proteins, nucleic acid molecules, antibodies, transformed cells and
XX expression vector are useful for detecting expression or levels of an
XX OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX cells in vitro. The antisense sequences and antibodies can be used to
XX promote growth of cells in vitro. Cell growth can be promoted by
XX interfering with the OGFR ligand-receptor system, especially where a
XX subject suffers from a tissue wound. Treating cancer comprises enhancing
XX the function of the OGFR ligand-receptor system in cancerous cells of a
XX patient or administering the OGFR nucleic acid to the patient.
XX
XX Sequence 2348 BP; 485 A; 738 C; 826 G; 299 T; 0 other;
XX
XX
XX alignment_scores:
XX Quality: 41.00 Length: 9
XX Ratio: 4.556 Gaps: 0
XX Percent Similarity: 100.000 Percent Identity: 77.778
XX
XX alignment_block:
XX US-09-444-281-27 x AAA28525/rev
XX
XX Align seg 1/1 to reverse of: AAA28525 from: 1 to: 2348
XX
XX 1 H1SGTUAAGIUPROGLUAGIUPRO 9

```

```

|||||
2165 CACGAAGCAGACCTGAGCGGAGTCCG 2139
seq_name: /SIDS8/gcgdata/geneseq/geneseq/NA2000.DAT:AAA28523
seq_documentation_block:
ID AAA28523 standard; cDNA; 2408 BP.
XX
XX AAA28523;
AC
XX 29-AUG-2000 (first entry)
XX
XX Human opioioid growth factor receptor cDNA spliced version 8.
XX
XX OGFR; opioioid growth factor receptor; growth inhibitor; proliferative;
XX cytostatic; vulnerary. gene therapy; antagonist; chromosome 20q13.3; ss.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX 5'UTR 1..33
XX FT /*tag= a
XX FT 34..2127
XX CDS /*tag= b
XX FT /product= Opioid_growth_factor_receptor
XX FT 2128..2408
XX FT /*tag= c
XX
XX 3'UTR
XX
XX WO200026340-A2.
XX
XX 11-MAY-2000.
XX
XX 02-NOV-1999; 99WO-US25802.
XX
XX 03-NOV-1998; 98US-0106879.
XX
XX (PENN-) PENN STATE RES FOUND.
XX
XX Zagon IS, McLaughlin PJ, Verderame MF;
XX WPI; 2000-365594/31.
XX DR P-PSDB; AAY92809.
XX
XX New cDNA encoding rat and human opioioid growth factor receptors which
XX modulate cell growth, useful for treating cancer
XX
XX Claim 1; Page 79-81; 91pp; English.
XX
XX Primers generated from rat opioioid growth factor receptor (OGFR) cDNA were
XX used to clone a fragment of the human OGFR cDNA. The complete sequence of
XX human OGFR was assembled with a combination of 3' and 5' RACE. 5' RACE
XX consistently yielded a single species of cDNA, while the 3' RACE revealed
XX extensive alternative splicing. The alternate splice forms were missing
XX the imperfect repeats or differed in the number of imperfect repeats. The
XX human OGFR gene chromosomal location was determined by FISH as 20q13.3.
XX expression vector are useful for detecting expression or levels of an
XX OGFR in a tissue. OGFR nucleic acids can be used to inhibit growth of
XX cells in vitro. The antisense sequences and antibodies can be used to
XX promote growth of cells in vitro. Cell growth can be promoted by
XX interfering with the OGFR ligand-receptor system, especially where a
XX subject suffers from a tissue wound. Treating cancer comprises enhancing
XX the function of the OGFR ligand-receptor system in cancerous cells of a
XX patient or administering the OGFR nucleic acid to the patient.
XX
XX Sequence 2408 BP; 500 A; 762 C; 844 G; 302 T; 0 other;
XX
XX
XX alignment_scores:
XX Quality: 41.00 Length: 9
XX Ratio: 4.556 Gaps: 0
XX Percent Similarity: 100.000 Percent Identity: 77.778
XX
XX alignment_block:

```

US-09-444-281-27 x AAA28523/rev ..

Align seg 1/1 to reverse of: AAA28523 from: 1 to: 2408

1 HisGLuAlaGluProGluAlaGluPro 9
|||||
2225 CACGAGCAGAGCCTGAGGAGGCCG 2199

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAH17441

seq_documentation_block:

ID AAH17441 standard; cDNA; 2409 BP.

AAH17441;

26-JUN-2001 (first entry)

Human cDNA sequence SEQ ID NO:16891.

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

Homo sapiens.

EP1074617-A2.

07-FEB-2001.

28-JUL-2000; 2000BP-0116126.

29-JUL-1999; 99JP-0248036.

27-AUG-1999; 99JP-0300253.

11-JAN-2000; 2000JP-018776.

02-MAY-2000; 2000JP-0183767.

09-JUN-2000; 2000JP-0241899.

(HELI-) HELIX RES INST.

Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

WPI; 2001-318749/34.

Claim 8; SEQ ID 16891; 2537BP + CD ROM; English.

The present invention describes primer sets for synthesizing 5602

full-length cDNAs defined in the specification. Where a primer set

comprises: (a) an oligo-dT primer and an oligonucleotide complementary

to the complementary strand of a polynucleotide which comprises one of

the 5602 nucleotide sequences defined in the specification, where the

oligonucleotide comprises at least 15 nucleotides; or (b) a combination

of an oligonucleotide comprising a sequence complementary to the

complementary strand of a polynucleotide which comprises a 5'-end

sequence and an oligonucleotide comprising a sequence complementary to a

polynucleotide which comprises a 3'-end sequence, where the

oligonucleotide comprises at least 15 nucleotides and the combination

of the 5'-end sequence/3'-end sequence is selected from those defined in

the specification. The primer sets can be used in antisense therapy and

particularly full-length cDNAs. The primers are also useful for the

detection and/or diagnosis of the abnormality of the proteins encoded by the

present invention.

alignment_scores:

Quality: 41.00

Ratio: 4.556

Percent Similarity: 100.000

Percent Identity: 77.778

alignment_block:

US-09-444-281-27 x AAH17441/rev ..

Align seg 1/1 to reverse of: AAH17441 from: 1 to: 2409

1 HisGLuAlaGluProGluAlaGluPro 9
|||||
2157 CACGAGCAGAGCCTGAGGAGGCCG 2131

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAH18239

seq_documentation_block:

ID AAH18239 standard; cDNA; 5796 BP.

AAH18239;

26-JUN-2001 (first entry)

Human cDNA sequence SEQ ID NO:18182.

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

Homo sapiens.

EP1074617-A2.

07-FEB-2001.

28-JUL-2000; 2000BP-0116126.

29-JUL-1999; 99JP-0248036.

27-AUG-1999; 99JP-0300253.

11-JAN-2000; 2000JP-018776.

02-MAY-2000; 2000JP-0183767.

09-JUN-2000; 2000JP-0241899.

(HELI-) HELIX RES INST.

Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

WPI; 2001-318749/34.

Claim 8; SEQ ID 18182; 2537BP + CD ROM; English.

The present invention describes primer sets for synthesizing 5602

full-length cDNAs defined in the specification. Where a primer set

comprises: (a) an oligo-dT primer and an oligonucleotide complementary

to the complementary strand of a polynucleotide which comprises one of

the 5602 nucleotide sequences defined in the specification, where the

oligonucleotide comprises at least 15 nucleotides; or (b) a combination

of an oligonucleotide comprising a sequence complementary to the

complementary strand of a polynucleotide which comprises a 5'-end

sequence and an oligonucleotide comprising a sequence complementary to a

polynucleotide which comprises a 3'-end sequence, where the

oligonucleotide comprises at least 15 nucleotides and the combination

of the 5'-end sequence/3'-end sequence is selected from those defined in

```

CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.
XX
SQ Sequence 5796 BP; 1434 A; 1616 C; 1531 G; 1215 T; 0 other;
XX
alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500
XX
alignment_block:
US-09-444-281-27 x AAH18239/rev
XX
Align seg 1/1 to reverse of: AAH18239 from: 1 to: 5796
XX
1 HisgLuAaGluProGluAaGlu 8
XX |||||||||*|||||||
XX 3514 CATGAGCTGAGCCTCAAGCTGAA 3491
XX
seq_name: /SIDS8/gcgdata/geneseq/geneseq/AA2001.DAT:AAH18247
XX
seq_documentation_block:
XX ID AAH18247 standard; cDNA: 6048 BP.
XX
XX AAH18247:
XX
XX 26-JUN-2001 (first entry)
XX
XX Human cDNA sequence SEQ ID NO:18198.
XX
XX Human; primer: detection; diagnosis; antisense therapy; gene therapy; ss
XX
XX Homo sapiens.
XX
XX EP1074617-A2.
XX
XX 07-FEB-2001.
XX
XX 28-JUL-2000; 2000EP-0116126.
XX
XX 29-JUL-1999; 99JP-0248036.
XX 27-AUG-1999; 99JP-0300253.
XX 11-JAN-2000; 2000JP-0118776.
XX 02-MAY-2000; 2000JP-0183767.
XX 09-JUN-2000; 2000JP-0241899.
XX
XX (HELI-) HELIX RES INST.
XX
XX Oka T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
XX Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX WPI: 2001-318749/34.
XX
XX Primer sets for synthesizing polynucleotides, particularly the 5602
XX full-length cDNAs defined in the specification, and for the detection
XX and/or diagnosis of the abnormality of the proteins encoded by the
XX full-length cDNAs -
XX
XX Claim 8; SEQ ID 18198; 2537bp + CD ROM; English.
XX
XX The present invention describes primer sets for synthesizing 5602
XX full-length cDNAs defined in the specification. Where a primer set
XX comprises: (a) an oligo-dT primer and an oligonucleotide complementary
XX to the complementary strand of a polynucleotide which comprises one of
XX the 5602 nucleotide sequences defined in the specification, where the
XX oligonucleotide comprises at least 15 nucleotides; or (b) a combination
XX of an oligonucleotide comprising a sequence complementary to the
XX complementary strand of a polynucleotide which comprises a 5'-end

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CC sequence and an oligonucleotide comprising a sequence complementary to a
CC polynucleotide which comprises a 3'-end sequence, where the
CC oligonucleotide comprises at least 15 nucleotides and the combination of
CC the 5'-end sequence/3'-end sequence is selected from those defined in
CC the specification. The primer sets can be used in antisense therapy and
CC in gene therapy. The primers are useful for synthesising polynucleotides
CC particularly full-length cDNAs. The primers are also useful for the
CC detection and/or diagnosis of the abnormality of the proteins encoded by
CC the full-length cDNAs. The primers allow obtaining of the full-length
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632
CC represent oligonucleotides, all of which are used in the exemplification
CC of the present invention.
XX
XX
S0 Sequence 6048 BP; 1488 A; 1693 C; 1604 G; 1263 T; 0 other:

Alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

Alignment_block:
US-09-444-281-27 x AAH18247/rev ..

Align seg 1/1 to reverse of: AAH18247 from: 1 to: 6048

1 HiscGuAlaGluProGluAlaGlu 8
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3767 CARGAGCTGAGCCTCAGGCTGAA 3744

seq_name: /SID58/gcgdata/geneseq/geneseqn/NA2001.DAT:AAI61181

seq_documentation_block:
ID AAI61181 standard; cDNA; 8605 BP.
AC AAI61181;
XX
XX 22-OCT-2001 (first entry)
DT
XX
DE Human polynucleotide SEQ ID NO 5170.
XX
XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
XX peripheral nervous system; neuropathy; central nervous system; CNS;
XX Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
XX amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
XX chemokine; thrombolytic; drug screening; arthritis; inflammation;
XX leukaemia; ss.
XX
XX Homo sapiens.
XX OS
XX WO200153312-A1.
XX
XX 26-JUL-2001.
XX
XX 26-DEC-2000; 2000WO-US34263.
XX
XX 21-JAN-2000; 2000US-0488725.
XX 25-APR-2000; 2000US-0552317.
XX 09-JUL-2000; 2000US-0598042.
XX 19-JUL-2000; 2000US-0620312.
XX 03-AUG-2000; 2000US-0653450.
XX 14-SEP-2000; 2000US-0662191.
XX 19-OCT-2000; 2000US-0693036.
XX 29-NOV-2000; 2000US-0727344.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
XX Wang J, Wang Z, Wehtman T, Xu C, Xue AJ, Yang Y, Zhang J;
XX Zhao QA, Zhou P, Goodrich R, Drmanac RT;

```

DR WPI; 2001-442253/47.
DR P-PSDB; AAM42025.
XX Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
XX
PS Claim 1: SEQ ID NO 5170; 10078bp; English.
XX
CC The invention relates to human nucleic acids (AA157798-AA161369) and
CC the encoded polypeptides (AAM38642-AAM42213) with neurotrophic,
CC immunosuppressant and cytoskeletal activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC localized neuropathies and central nervous system diseases, such as
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
CC utilization of the activities such as: Immune system suppression,
CC activation/inhibition activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
XX
SQ Sequence 8605 BP; 2079 A; 2209 C; 2192 G; 2125 T; 0 other;
XX
alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500
XX
alignment_block:
US-09-444-281-27 x AA161181 ..
XX
Align seg 1/1 to: AA161181 from: 1 to: 8605
XX
1 HisGUAAGAGUProGUAGUAGU 8
|||||
5995 CATGAGCTGAGCCTCAGCGTGAA 6018
XX
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA2001.DAT:AA159395
XX
seq_documentation_block:
ID AA159395 standard; cDNA; 8840 BP.
XX
AC AA159395;
XX
DT 22-OCT-2001 (first entry)
XX
DE Human polynucleotide SEQ ID NO 1598.
XX
KW Human; neurotrophic; immunosuppressant; cytoskeletal; gene therapy; cancer;
KW peripheral nervous system; neuropathy; central nervous system; CNS;
KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
KW leukaemia; ss.
XX
OS Homo sapiens.
XX
PN WO20015312-A1.
XX
PD 26-JUL-2001.
XX
PE 26-DEC-2000; 2000WO-US34263.
XX
PR 21-JAN-2000; 2000US-0488725.
PR 25-APR-2000; 2000US-0552317.
PR 09-JUL-2000; 2000US-0588042.
PR 19-JUL-2000; 2000US-0620312.
PR 03-AUG-2000; 2000US-0653450.

PR 14-SEP-2000; 2000US-0662191.
PR 19-OCT-2000; 2000US-0693036.
PR 29-NOV-2000; 2000US-0727344.
XX
XX (HYSE-) HYSEQ INC.
XX
PI Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
PI Zhao QA, Zhou P, Goodrich R, Drmanac RT;
XX
DR WPI; 2001-442253/47.
DR P-PSDB; AAM40239.
XX
PT Novel nucleic acids and polypeptides, useful for treating disorders
PT such as central nervous system injuries -
XX
PS Claim 1: SEQ ID NO 1598; 10078bp; English.
XX
CC The invention relates to human nucleic acids (AA157798-AA161369) and
CC the encoded polypeptides (AAM38642-AAM42213) with neurotrophic,
CC immunosuppressant and cytoskeletal activity. The polynucleotides are useful
CC in gene therapy. A composition containing a polypeptide or polynucleotide
CC of the invention may be used to treat diseases of the peripheral nervous
CC system, such as peripheral nervous injuries, peripheral neuropathy and
CC localized neuropathies and central nervous system diseases, such as
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
CC utilization of the activities such as: Immune system suppression,
CC activation/inhibition activity, chemotactic/chemokinetic activity, haemostatic
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
CC C.N.S disorders.
CC Note: The sequence data for this patent did not form part of the printed
CC specification.
XX
SQ Sequence 8840 BP; 2118 A; 2331 C; 2255 G; 2136 T; 0 other;
XX
alignment_scores:
Quality: 40.00 Length: 8
Ratio: 5.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500
XX
alignment_block:
US-09-444-281-27 x AA159395/rev ..
XX
Align seg 1/1 to reverse of: AA159395 from: 1 to: 8840
XX
1 HisGUAAGAGUProGUAGUAGU 8
|||||
3604 CATGAGCTGAGCCTCAGCGTGAA 3581
XX
seq_name: /SIDS8/gcgdata/geneseq/geneseqn/NA1998.DAT:AAV68396
XX
seq_documentation_block:
ID AAV68396 standard; cDNA to mRNA; 9408 BP.
XX
AC AAV68396;
XX
DT 05-MAY-1999 (first entry)
XX
DE Human BA22-alpha cDNA.
XX
KW Transcriptional regulator; transcription; BA21-alpha; bromodomain; BA2;
KW atypical zinc finger; testis; human; tumour; BA21-beta; BA22-alpha; drug;
KW BA22-beta; treatment; cancer; proliferative disorder; screening; ds.
XX
OS Homo sapiens.
XX
PH Key Location/Qualifiers
FH CDS 740..6376
FT /*tag= a
FT /product= "BA22-alpha"

XX WO9847920-A1.
 PN
 XX
 PD 29-OCT-1998.
 XX
 PF 17-APR-1998; 98WO-JP01783.
 XX
 PR 24-OCT-1997; 97JP-0310027.
 PR 18-APR-1997; 97JP-0116570.
 XX
 PA (CHUG-) CHUGAI RES INST MOLECULAR MEDICINE INC.
 XX
 PI Jones MH;
 XX
 DR WPI: 1998-583603/49.
 DR P-PSDB; AAW81170.
 XX
 PT Transcriptional regulator gene family containing bromodomain - may
 PT be expressed in testis tissue and is useful for treatment of cancer
 PT and other proliferative disorders
 XX
 PS Claim 2; Page 72-88; 187pp; Japanese.
 XX
 CC This sequence encodes the human BA22-alpha protein, a member of a
 CC family of transcriptional regulator genes containing a bromodomain (BA2,
 CC Bromodomain with Atypical zinc finger) which are expressed specifically
 CC in testis tissue and also in certain tumour lines. Transgenic cells may
 CC be used for the preparation of the BA21-alpha, BA21-beta, BA22-alpha and
 CC BA22-beta proteins. These proteins can be used in the treatment of cancer
 CC and other proliferative disorders, and in screening of compounds for
 CC their binding ability to the expression products (e.g. for use as drugs
 CC by modulation of transcriptional regulation).
 XX
 SQ Sequence 9408 BP; 2279 A; 2458 C; 2383 G; 2280 T; 8 other;

alignment_scores:
 Quality: 40.00 Length: 8
 Ratio: 5.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
 US-09-444-281-27 x AAV68396/rev ..
 Align seg 1/1 to reverse of: AAV68396 from: 1 to: 9408
 1 HtSGTuaIaGluProGluAlaGlu 8
 ||||||||||||||||:||||||
 4343 CATGAAGCTGAGCTCAGAGTGAA 4320

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REFERENCE	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Nematocera; Culicoidae; Anopheles.
AUTHORS	1 (bases 1 to 848)
TITLE	Genoscope.
JOURNAL	Direct Submission
REFERENCE	Submitted (16-FEB-2000) Genoscope - Centre National de Sequencage
AUTHORS	Bp 191 91006 EVR cedex - FRANCE (E-mail : segreif@genoscope.cns.fr)
TITLE	2 (bases 1 to 848)
JOURNAL	Roeh,C.W., Brey,P.T., Ke,Z., Collins,F.H. and Weissenbach,J.
COMMENT	Submitted (16-FEB-2000) BMML Institut Pasteur, 25, rue du Dr. Roux, Paris 75015, France
FEATURES	This clone is from an A. gambiae BAC library provided by F.H. Collins and sequenced by Genoscope in collaboration with the Laboratory of Biochem. and Biol. Molec. of Insects, Institut Pasteur.
source	location/Qualifiers
	1..848
	/organism="Anopheles gambiae"
	/strain="PE8"
	/db_xref="taxon:7165"
	/clone="18M20"
	/clone_lib="Notredame1"
	/note="end : 17"
BASE COUNT	216 a 157 c 204 g 248 t 23 others
ORIGIN	
alignment_scores:	
Quality:	45.00 Length: 11
Ratio:	4.500 Gaps: 0
Percent Similarity:	90.909 Percent Identity: 72.727
alignment_block:	
US-09-444-281-27 x CNS01L90/rev ..	
Align seg 1/1 to reverse of: CNS01L90 from: 1 to: 848	
1 HISG1UALAG1UPROG1UALAG1PRO1Lmet 11	
::: ::: :::	
666 CAGGATCCATACCGATCGAATCGAACCTATCATG 634	
seq_name: gb-gss:A2208017	
seq_documentation_block:	
LOCUS A2208017 923 bp DNA GSS 31-AUG-2000	
DEFINITION SP_0134.B1.D06.T7A Strongylocentrotus purpuratus, purple sea urchin	
clone, sperm genomic BAC library Strongylocentrotus purpuratus genomic	
clone plate=134 Col=11 Row=H, DNA sequence.	
ACCESSION A2208017	
VERSION A2208017.1 GI:8420201	
KEYWORDS GSS.	
SOURCE Strongylocentrotus purpuratus.	
ORGANISM Strongylocentrotus purpuratus	
REFERENCE Eukaryota; Metazoa; Echinodermata; Eleutherozoa; Echinozoa;	
AUTHORS Echinoidae; Euechinoidae; Echinacea; Echinoida;	
REFERENCE Strongylocentrotidae; Strongylocentrotus.	
1 (bases 1 to 923)	
Swartzell,S., Wallace,J.C., Pouska,A.J., Livingston,B.T., Wray	
G.A., Ettensohn,C.A., Lehrach,H., Britten,R.J., Davidson,E.H. and	
Hood,L.	
A sea urchin genome project: Sequence scan, virtual map, and	
additional resources	
Proc. Natl. Acad. Sci. U. S. A. 97 (17), 9514-9518 (2000)	
20402566	
COMMENT Contact: Cameron, RA, Davidson, EH, Hood, L	
MEDLINE Division of Biology 156-29	
California Institute of Technology	
Pasadena California 91125, USA	

Fax: (626) 793-3047
Email: acamerone@caltech.edu
Plate: 134 row: 18 column: 11
Seq primer: T7
Class: BAC ends
High quality sequence stop: 923.
Location/Qualifiers

FEATURES
source

1. 923
/organism="Strongylocentrotus purpuratus"
/db_xref="taxon:7668"
/clone="plate-134 Col-11 Row-H"
/clone_lib="Strongylocentrotus purpuratus, purple sea
urchin, sperm genomic BAC library"
/note="Organ: sperm; Vector: BAC3.6; BAC clones in E-Coli
DH10B"

BASE COUNT 173 a 216 c 203 g 331 t
ORIGIN

alignment_scores:

Quality: 45.00 Length: 9
Ratio: 5.625 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-09-444-281-27 x AZ208017/rev

Align seg 1/1 to reverse of: AZ208017 from: 1 to: 923

1 HsGluAlaGluProGluAlaGluPro 9
|||||
55 CACGAAGCCAGCCAGCCAGCACCA 29

seq_name: gb_est2:BF525048

seq_documentation_block:

LOCUS BF525048 343 bp mRNA EST 11-DEC-2000
DEFINITION UI-R-AD0-vz-d-06-0-UI-r1 UI-R-AD0 Rattus norvegicus cDNA clone
UI-R-AD0-vz-d-06-0-UI 5', mRNA sequence.

ACCESSION BF525048
VERSION BF525048.1 GI:11633015

KEYWORDS

SOURCE

ORGANISM

Norman rat.
Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 343)

Bonaldo, M.F., Lennon, G. and Soares, M.B.
Normalization and subtraction: two approaches to facilitate gene
discovery

Genome Res. 6 (9), 791-806 (1996)

JOURNAL
MEDLINE
COMMENT

Contact: Soares, MB.
Program for Rat Gene Discovery and Mapping
University of Iowa
451 Eckstein Medical Research Building Iowa City, IA 52242, USA
Tel: 319 335 8250
Fax: 319 335 9565
Email: msoares@blue.weeg.uiowa.edu
cDNA Library Preparation: M.B. Soares Lab Clone distribution:
Clones will be available through Research Genetics (www.resgen.com)
This clone is also available through the I.M.A.G.E. Consortium at
LNL (info@image.lnl.gov). IMAGE ID= 1794853
Seq primer: M13 Forward.

FEATURES

source

1. 343
/organism="Rattus norvegicus"
/strain="Sprague-Dawley"
/db_xref="taxon:10116"
/clone="UI-R-AD0-vz-d-06-0-UI"
/clone_lib="UI-R-AD0"
/dev_stage="adult"

/lab_host="DH10B (Life Technologies)"
/note="Vector: pTZ19D-Pac (Pharmacia) with a modified
polylinker. Site 1: Not I; Site 2: Eco RI; The UI-R-AD0
library is a non-normalized library constructed from 15
dpc rat atrium. The tag is a string of 5 nucleotides
present between the Not I site and the oligo-dT track.
The library was constructed as described by Bonaldo,
Lennon and Soares, Genome Research 6: 791-806, 1996.
Tissue provided by Jim Lin, Department of Biology,
University of Iowa."

BASE COUNT 106 a 83 c 95 g 59 t
ORIGIN

alignment_scores:

Quality: 43.00 Length: 11
Ratio: 3.909 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 63.636

alignment_block:

US-09-444-281-27 x BF525048

Align seg 1/1 to: BF525048 from: 1 to: 343

1 HsGluAlaGluProGluAlaGluProIleMet 11
|||||
127 CATCATGCAGACGCTCATGCAGACCTCTTGC 159

seq_name: gb_est1:AW140836

seq_documentation_block:

LOCUS AW140836 552 bp mRNA EST 30-OCT-1999
DEFINITION EST290918 Normalized rat heart, Bento Soares Rattus sp. cDNA clone
RG1AV22 5' end similar to peroxisomal farnesylated protein, mRNA
sequence.

ACCESSION AW140836
VERSION AW140836.1 GI:6160674

KEYWORDS

SOURCE

ORGANISM

Rattus sp.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

1 (bases 1 to 552)

Lee, N.H., Glodek, A., Chandra, I., Mason, T.M., Quackenbush, J.,
Kerlavage, A.R. and Adams, M.D.
Rat Genome Project: Generation of a Rat EST (RESt) Catalog & Rat
Gene Index

Unpublished (1998)
Other ESTs: TC85992
Contact: Lee, NH
The Institute for Genomic Research
9712, Medical Center Drive, Rockville, MD 20850, USA
Tel: (301)-838-3529
Fax: (301)-838-0208
Email: nhlee@tigr.org

For clone availability, additional sequence and expression
information related to this EST please check the TIGR Rat Gene
Index (<http://www.tigr.org/tdb/t91/t91.html>). To order a clone
contact the ATCC (<http://www.atcc.org/atcc.html>).
Seq primer: M13 Reverse.

FEATURES

source

1. 552
/organism="Rattus sp."
/db_xref="taxon:10118"
/clone="RG1AV22"
/clone_lib="Normalized rat heart, Bento Soares"
/note="Organ: heart; Vector: pTZ19D-Pac; Site 1: EcoRI;
Site 2: NotI"
BASE COUNT 155 a 138 c 149 g 110 t
ORIGIN


```

template preparation) comprising: a) a set of about 1,000 arrayed clones from each of the five non-normalized libraries of brain (CTOs), heart (CSOs), kidney (CKOs), aorta (CWOs), and placenta (CXOs). The resulting pool of approximately 5,000 clones represented about 33.3% of the final driver population. A set of about 2,000 arrayed clones from each of the five normalized libraries of brain (CTO), heart (CSO), kidney (CKO), aorta (CWO), and placenta (CXO). The resulting pool of about 10,000 clones represented about 66.6% of the final driver population.
TAG_Library=UI-R-DKO
TAG_Tissue=rat heart pool
TAG_SEQ=AFAAGATTAAC"
BASE COUNT      126 a          179 c          169 g          217 t
ORIGIN

alignment_scores:
    Quality:      43.00           Length:       11
    Ratio:        3..909           Gaps:         0
Percent Similarity: 100.000     Percent Identity: 63.636

alignment_block:
US-09-444-281-27 x BI295006/rev ..

Align seg 1/1 to reverse of: BI295006 from: 1. to: 691

1 HHSGLUAGLUPROGSLUAGLUPPOILEMET 11
|||||:|||||:|||||:|||||:|||||:
435 CATCATGCAGAGCGCTCATGCAGACTCTTGTC 403

seq_name: gb_est2.BF259969

seq_documentation_block:
LOCUS      BF259969              836 bp            mRNA                    EST                23-FEB-2001
DEFINITION HVSMET0020M02f Hordeum vulgare seedling root EST library HVCDNA0007
            (etiolated and unstressed) Hordeum vulgare cDNA clone
ACCSSION   BF259969
VERSION    BF259969.2 GI:13120375
KEYWORDS   EST.
SOURCE     barley.
ORGANISM   Hordeum vulgare
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooideae
            ; Triticeae; Hordeum.
REFERENCE   1 (bases 1 to 836)
AUTHORS    Wang,R., Close,T.J., Kleinhofs,A., Wise,R., Begum,D., Fritsch,D., Yu
            ,Y., Anderson,H., Dale,J., Henry,D., Kennole,S., Palmer,M., Rambo
            ,T., Sakri,C., Schwartzbeck,T., Simmons,J., Choi,D.W., Main,D. and
            Wood,T.
FEATURES             (1)..(836)
     TITLE               Development of a genetically and physically anchored EST resource
                        for barley genomics
     JOURNAL              unpublished (2000)
     COMMENT              On Nov 16, 2000 this sequence version replaced gi:1189082.
                        Contact: Wing RA
                        Clemson University Genomics Institute
                        Clemson University
                        100 Jordan Hall, Clemson, SC 29634, USA
                        Tel.: 864 656 7288
                        Fax: 864 656 4293
                        Email: rwing@clemson.edu
     SEG PRIMER          AATTACCCGTCACTAAGGG
                        High quality sequence stop: 512.
                        Location/Organism
                        1..836
                        /organism="Hordeum vulgare"
                        /cultivar="Morex"
                        /db_xref="taxon:4513"
                        /clone="HVSMET0020M02f"
                        /clone_lib="Hordeum vulgare seedling root EST library
                        HVCDNA0007 (etiolated and unstressed)"
                        /tissue_type="Seedling root"

```

```

/lab host="TJc121"
/note="vector: lambdaZAP; Site_1: EcoRI; Site_2: XhoI. For
more details on library preparation and sequence analysis
see http://www.genome.clemson.edu/projects/bailey/ To
order a clone see http://www.genome.clemson.edu/orders"
BASE COUNT      163 a      233 c      275 g      165 t
ORIGIN

alignment_scores:
    Quality:      43.00          Length:      9
    Ratio:        4:.778         Gaps:      0
Percent Similarity: 100.000     Percent Identity: 77.778

alignment_block:
US-09-444-281-27 x BF259969 ..

Align seg 1/1 to: BF259969 from: 1 to: 836

1 HISGLUAGLUPROGUAAGLUPRO 9
||||| ||||| ||||| ||||| |||||
614 CATGATCGCTGAACGGAAAGCTGAGCCG 640

seq_name: gb_est1:BE037128

seq_documentation_block:
LOCUS       BE037128      1247 bp      mRNA           EST       07-JUN-2000
DEFINITION  MP15E02 MP Mesembryanthemum crystallinum cDNA 5' similar to
auxin-regulated protein, mRNA sequence.
ACCESSION   BE037128
VERSION     BE037128.1 GI:8332144
KEYWORDS    EST.
SOURCE      common ice plant.
ORGANISM    Mesembryanthemum crystallinum
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophytes; Magnoliophyta; eudicotyledons; core eudicots;
            Caryophyllidae; Caryophyllales; Alzooceae; Mesembryanthemum.
REFERENCE   1 (bases 1 to 1247)
AUTHORS    Bonnett,H.J., Borchert,C., Brazille,S., Brooks,J., Eaton,M., Ferreira
            ,H., Kawasak,S., McCollough,A., Michalowski,C.B., Palacio,C.,
            Scara,G., Wheeler,M. and Zepeda,G.R.
COMMENT     Functional Genomics of Plant Stress Tolerance
            Unpublished (2000)
            Contact: Michalowski,C.B.
            University of Arizona
            Bio Sciences West room 513, Tucson, AZ 85721, USA
            Tel: 520-621-7982
            Fax: 520-621-1697
            Email: cbm@u.arizona.edu.

FEATURES             Location/Qualifiers
     source            1..1247
                     /organism="Mesembryanthemum crystallinum"
                     /db_xref="taxon:3544"
                     /clone_lib="MP"
                     /isolate_type="apical meristem and leaf primordia"
                     /disease="6 weeks"
                     /note="3 d 500mm NaCl"
BASE COUNT      270 a      300 c      242 g      301 t      134 others
ORIGIN

alignment_scores:
    Quality:      43.00          Length:      10
    Ratio:        5:.375         Gaps:      0
Percent Similarity: 80.000     Percent Identity: 70.000

alignment_block:
US-09-444-281-27 x BE037128/rev ..

Align seg 1/1 to reverse of: BE037128 from: 1 to: 1247

||||| ||||| ||||| ||||| |||||
1 HISGLUAGLUPROGUAAGLUPROIE 10
||||| ||||| ||||| ||||| |||||

```

1000

Align seg 1/1 to reverse of: T70690 from: 1 to: 323

1 H18GUALAGUPROGLUAGLUPRO 10
||||| ||||||| |||||||
36 CACGACCAAGAACCTGAGACGACGAGCTG 7

seq_name: gb_est1:BI240337

seq_documentation_block:

LOCUS BI240337 393 bp mRNA EST 12-JUL-2001
DEFINITION R337162.5prime RE Drosophila melanogaster normalized Embryo p1c-1
Drosophila melanogaster cDNA clone R337162.5 similar to CG7611:
Faan0007611 located on: 3L 78B1-78B1; 05/12/2001, mRNA sequence.

ACCESSION BI240337
VERSION BI240337.1 GI:14708940
KEYWORDS EST
SOURCE fruit fly.

ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 (bases 1 to 393)
Stapleton, M., Brokstein, P., Hong, L., Tyler, D., Berman, B., Carlson
, J., Champe, M., Chavez, C., Dorsett, V., Farfan, D., Frise, E., George
, R., Gonzalez, M., Guarin, H., Harris, N., Li, P., Liao, G., Misra, S.,
Mungall, C. J., Nunoo, J., Pacleb, J., Paragas, V., Park, S.,
Phonsavong, S., Wan, K., Yu, C., Lewis, S. E., Celniker, S. and Rubin
, G. M.

TITLE BDGP/HMM RE Drosophila EST Project
JOURNAL Unpublished (2001)
COMMENT Contact: Stapleton, M.

BASE COUNT 118 a 86 c 89 g 126 t
ORIGIN

FEATURES
source
1. 393
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/clone="RE37162"
/clone_1lb="RE Drosophila melanogaster normalized Embryo
p1c-1"
/sex="male and female"
/dev_stage="0-24 hours mixed stage embryonic"
/lab_host="DH5-alpha Tona"
/note="Organ: embryo; Vector: pPIC1; Site_1: XhoI; Site_2:
BamHI; Library was kindly generated by Piero Carninci at
the RIKEN. The library was normalized and excised using
Cre recombinase. Plasmid cDNA library."

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x BI240337
Align seg 1/1 to: BI240337 from: 1 to: 393

2 GUUAGLUPROGLUAGLUPRO 9
||||| ||||||| |||||||
190 GAGGCTGAGCCTGAGGACGAGCT 213
seq_name: gb_est1:AU052906

seq_documentation_block:

LOCUS AU052906 419 bp mRNA EST 28-APR-1999
DEFINITION AU052906 Dictyostelium discoideum SL (H. Urushihara) Dictyostelium
discoideum cDNA clone SLF303, mRNA sequence.

ACCESSION AU052906
VERSION AU052906.1 GI:4701389
KEYWORDS EST

SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum.
Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.

REFERENCE 1 (bases 1 to 419)
Morio, T., Urushihara, H., Saito, T., Ugawa, Y., Mizuno, H., Yoshida, M.,
Yoshino, R., Mitra, B. N., Pl, M., Saito, T., Takemoto, K., Yasukawa, H.,
Williams, J., Meeda, M., Takeuchi, I., Ochiai, H. and Tanaka, Y.

TITLE Developmental cDNA in Dictyostelium discoideum
JOURNAL Unpublished (1998)
COMMENT Contact: Hideko Urushihara
Institute of Biological Sciences
University of Tsukuba
3-3-10 Ten-nodai, Tsukuba, Ibaraki 305, Japan
Email: d402huesakura.cc.tsukuba.ac.jp
PROJECT = Dictyostelium discoideum cDNA project in Japan.

FEATURES
source
1. 419
/organism="Dictyostelium discoideum"
/strain="AX4"
/db_xref="taxon:44689"
/clone="SLF303"
/clone_1lb="Dictyostelium discoideum SL (H. Urushihara)"
/dev_stage="slug"

BASE COUNT 118 a 86 c 89 g 126 t
ORIGIN

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x AU052906/rev
Align seg 1/1 to reverse of: AU052906 from: 1 to: 419

2 GUUAGLUPROGLUAGLUPRO 9
||||| ||||||| |||||||
164 GAGGCGGAGACCTGAGCGGAGCT 141
seq_name: gb_est1:AU923872

seq_documentation_block:
LOCUS AU923872 435 bp mRNA EST 19-JUL-2000
DEFINITION WS1_30-B09.D1_A002 Water-stressed 1 (WS1) Sorghum bicolor cDNA,
mRNA sequence.

ACCESSION AU923872
VERSION AU923872.1 GI:8089697
KEYWORDS EST
SOURCE sorghum.

ORGANISM sorghum bicolor
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACC
clade; Panicoidae; Andropogoneae; Sorghum.

REFERENCE 1 (bases 1 to 435)
Cordonnier-Pratt, M.-M., Gingle, A., Marsala, C., Sudman, M. and Pratt
, L. H.

TITLE An EST database from Sorghum: water-stressed plants
JOURNAL Unpublished (2000)
COMMENT Contact: Cordonnier-Pratt MM
Department of Botany
The University of Georgia
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860

Fax: 706 542 1805
Email: mmpratt@uga.edu
Sequences have been trimmed to exclude POLYA, vector and regions
below Phred quality 16. The threshold for highest quality sequence
is 20.

Seq primer: JEN REV
High quality sequence stop: 368
POLYA-No.

FEATURES
source

1.435 location/Qualifiers
/organism="Sorghum bicolor"
/db_xref="taxon:4558"
/clone_lib="Water-stressed 1 (WS1)"
/note="Organ: Mix of 5-week old plants on days 7 & 8 after
water was withheld; Vector: Lambda Zap; Site_1: XhoI;
Site_2: EcoRI; The library was made from poly-A RNA in the
cloning vector Lambda Zap II. Clones to be sequenced were
prepared by mass excision."
156 a 162 g 84 g 33 t
BASE COUNT
ORIGIN

alignment_scores:
Quality: 42.00 Length: 10
Ratio: 4.667 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-27 x AW923872

Align seg 1/1 to: AW923872 from: 1 to: 435

1 HisqluAlagluProgluAlagluProfile 10
|||||
2 CACGAGGCTGAGCCTCAACCTAAACCATTA 31

GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:40:32 ; Search time 50.17 Seconds
(Without alignments)
32.071 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAPEAPEPIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues

Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL_17:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted, by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	42	71.2	880	5	017338 caenorhabd
2	42	71.2	885	5	09TXR9 caenorhabd
3	42	71.2	930	5	017339 caenorhabd
4	41	69.5	874	5	09V010 caenorhabd
5	40	67.8	248	12	065200 drosophila
6	40	67.8	1380	5	09V108 african swi
7	39	66.1	251	3	09US24 schizosacch
8	39	66.1	304	5	09NAB8 caenorhabd
9	39	66.1	340	5	09NAB2 caenorhabd
10	39	66.1	408	5	096979 parophrys
11	39	66.1	417	5	09W4Y3 drosophila
12	39	66.1	435	5	09NEF8 drosophila
13	39	66.1	848	5	021489 caenorhabd
14	39	66.1	1192	5	09W475 drosophila
15	39	66.1	1277	12	008547 cercopithec
16	39	66.1	1279	12	066031 cercopithec
17	39	66.1	1458	3	09HE72 neotropora
18	39	66.1	1593	5	020207 caenorhabd
19	39	66.1	6797	2	09X993 streptomyce

20	38	64.4	150	1	09YFA9 aeropyrum p
21	38	64.4	200	4	09UD8 homo sapien
22	38	64.4	288	3	09P3H2 neotropora
23	38	64.4	592	3	059900 cyproccoc
24	38	64.4	682	2	09A808 caulobacter
25	38	64.4	1083	2	086637 streptomyce
26	37	62.7	20	4	09UD25 homo sapien
27	37	62.7	101	6	095284 sus scrofa
28	37	62.7	125	2	09WX24 streptomyce
29	37	62.7	175	10	09SYE4 arabidopsis
30	37	62.7	208	4	060937 homo sapien
31	37	62.7	248	11	063224 ratius norv
32	37	62.7	251	6	09GLN2 bos taurus
33	37	62.7	265	37	09JZP0 neisseria m
34	37	62.7	265	2	09JUS3 neisseria m
35	37	62.7	338	10	09M4K6 anthoceros
36	37	62.7	356	10	09M4K5 anthoceros
37	37	62.7	387	5	09VZC7 arabidopsis
38	37	62.7	401	5	09VZC6 drosophila
39	37	62.7	407	5	09NCB6 drosophila
40	37	62.7	439	11	09JMF5 mus musculu
41	37	62.7	439	11	09JMF5 mus musculu
42	37	62.7	439	11	09JMF5 mus musculu
43	37	62.7	583	4	09UKV2 homo sapien
44	37	62.7	612	5	09U003 caenorhabd
45	37	62.7	614	4	09UKV1 homo sapien

ALIGNMENTS

RESULT	1	PRELIMINARY	PRT	880 AA.
ID	017338			
AC	017338			
DT	01-JAN-1998 (TREMBLrel. 05, Created)			
DT	01-JAN-1998 (TREMBLrel. 05, Last sequence update)			
DT	01-NOV-1998 (TREMBLrel. 08, Last annotation update)			
DE	T23E7.2B PROTEIN.			
GN	T23E7.2B.			
OS	Caenorhabditis elegans.			
OC	Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;			
OC	Rhabditidae; Peloderinae; Caenorhabditis.			
OX	NCBI_TaxID=6239;			
RN	[1]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RX	MEDLINE=94150718; PubMed=7906398;			
RA	Wilson R., Ainscough R., Anderson K., Baynes C., Berks M.,			
RA	Bonfield J., Burton J., Connell M., Copesey T., Cooper J., Coulson A.,			
RA	Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,			
RA	Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,			
RA	Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,			
RA	Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,			
RA	Parsons J., Percy C., Rifken L., Roopra A., Saunders D., Showkeen R.,			
RA	Smaldon N., Smith A., Sonhammer E., Staden R., Sulston J.,			
RA	Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K., Waterston R.,			
RA	Watson A., Weinstock L., Wilkinson-Sprout J., Wohlman P.,			
RT	"2.2 Mb of contiguous nucleotide sequence from Chromosome III of C.			
RT	elegans.";			
RL	Nature 368:32-38(1994).			
RN	[2]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RA	Latreille P., Steillies L., Elliott G.,			
RL	Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.			
RN	[3]			
RC	SEQUENCE FROM N.A.			
RC	STRAIN-BRISTOL N2;			
RA	Waterston R.,			
RL	Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.			
DR	EMBL; AF026205; AAB71258.1;			
SO	SEQUENCE 880 AA; 95398 MW; 97A8A101E8FBA1C1 CRC64;			

RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Idegam C.,
RA Jalali M., Kalush F., Karpen G.H., Ka.Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Krafc C., Kravitz S., Kuip D., Lai Z.,
RA Laslo P., Lai Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Matel B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merklov G., Mishina N.V., Mobarly C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
RA Palazzo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Slier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Sytkas R., Teclor C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weisenbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yen R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
DR EMBL: AEO03581: AAF51156.1:-
DR FlyBase: FBgn0031496; CG17258.
SQ SEQUENCE 874 AA: 103694 MW: 5F56DJCE7A01D9A CRC64;

Query Match 69.5% Score 41; DB 5; Length 874;
Best Local Similarity 70.0% Pred. NO. 49;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 2 EAEPEPEPIM 11
|||||
Db 376 EAEPEPEPPL 385

RESULT 5
ID 065200 PRELIMINARY; PRT: 248 AA.
AC 065200:
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE PE248R.
GN E248R.
OS African swine fever virus (ASFV).
OC Viruses; dsDNA viruses, no RNA stage; Asfarviridae;
OC African swine fever-like viruses.
OX NCBI_TaxID=10497;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=96036500; PubMed=7483270;
RA Yanez R.J., Rodriguez J.M., Nogal M.L., Yuste L., Enriquez C.,
RA Rodriguez J.F., Vinuela E.;
RT "Immune protection conferred by the baculovirus-related glycoprotein
RT of Hogdoto virus (Orthomyxoviridae).";
RL Virology 208:249-278(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=94233765; PubMed=8178480;
RA La Vega I., Gonzalez A., Blasco R., Calvo V., Vinuela E.;
RT "Nucleotide sequence and variability of the inverted terminal
RT repetitions of African swine fever virus DNA.";
RL Virology 201:152-156(1994).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=90219205; PubMed=2325203;
RA Gonzalez A., Calvo V., Almazan F., Almendral J.M., Ramirez J.C.,
RA La Vega I., Blasco R., Vinuela E.;
RT "Multigene families in African swine fever virus: family 360.";
RL J. Virol. 64:2073-2081(1990).

RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=90219204; PubMed=2325202;
RA Almendral J.M., Almazan F., Blasco R., Vinuela E.;
RT "Multigene families in African swine fever virus: family 110.";
RL J. Virol. 64:2064-2072(1990).
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=91134988; PubMed=1994575;
RA Camacho A., Vinuela E.;
RT "Protein p22 of African swine fever virus: an early structural protein
RT that is incorporated into the membrane of infected cells.";
RL Virology 181:251-257(1991).
RN [6]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RA Almazan F., Murguia J.R., Rodriguez J.M., La Vega I., Vinuela E.;
RL J. Gen. Virol. 0:0-0(0).
RN [7]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=94187118; PubMed=8139051;
RA Rodriguez J.M., Yanez R.J., Pan R., Rodriguez J.F., Salas M.L.,
RA Vinuela E.;
RT "Multigene families in African swine fever virus: family 505.";
RL J. Virol. 68:2746-2751(1994).
RN [8]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=93346971; PubMed=8393914;
RA Yanez R.J., Rodriguez J.M., Rodriguez J.F., Salas M.L., Vinuela E.;
RT "African swine fever virus thymidylate kinase gene: sequence and
RT transcriptional mapping.";
RL J. Gen. Virol. 74:1633-1638(1993).
RN [9]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=94065656; PubMed=8245848;
RA Alcant A., Angulo A., Vinuela E.;
RT "Mapping and sequence of the gene encoding the African swine fever
RT virion protein of M(r) 11500.";
RL J. Gen. Virol. 74:2317-2324(1993).
RN [10]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=93277388; PubMed=8503790;
RA Munoz M., Freije J.M., Salas M.L., Vinuela E., Lopez-Otin C.;
RT "Structure and expression in E. coli of the gene coding for protein
RT p10 of African swine fever virus.";
RL Arch. Virol. 130:93-107(1993).
RN [11]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=90357780; PubMed=2389555;
RA Blasco R., Lopez-Otin C., Munoz M., Bockamp E.O., Simon-Mateo C.,
RA Vinuela E.;
RT "Sequence and evolutionary relationships of African swine fever virus
RT thymidine kinase.";
RL Virology 178:301-304(1990).
RN [12]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;
RX MEDLINE=93281390; PubMed=8506138;
RA Yanez R.J., Boursnell M., Nogal M.L., Yuste L., Vinuela E.;
RT "African swine fever virus encodes two genes which share significant
RT homology with the two largest subunits of DNA-dependent RNA
RT polymerases.";
RL Nucleic Acids Res. 21:2423-2427(1993).
RN [13]
RP SEQUENCE FROM N.A.
RC STRAIN=BA71V;

RX MEDLINE-93353606; PubMed-8102411;
 RA Rodriguez J.M., Yanez R.J., Almazan F., Vinuela E., Rodriguez J.F.;
 RT "African swine fever virus encodes a CD2 homolog responsible for the
 RL adhesion of erythrocytes to infected cells."; J. Virol. 67:5312-5320(1993).
 RN [14]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94085774; PubMed-8262374;
 RA Yanez R.J., Rodriguez J.M., Bournell M., Rodriguez J.F., Vinuela E.;
 RT "Two putative African swine fever virus helicases similar to yeast
 RL and DExH." pre-mRNA processing proteins and vaccinia virus ATPases DILL
 Gene 134:161-174(1993).
 RN [15]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-90223993; PubMed-2327074;
 RA Lopez-Otin C., Freije J.M., Parra F., Mendez E., Vinuela E.;
 RT "Mapping and sequence of the gene coding for protein p72, the major
 RL capsid protein of African swine fever virus."; Virology 175:477-484(1990).
 RN [16]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94123986; PubMed-8293992;
 RA Rodriguez J.M., Yanez R.J., Rodriguez J.F., Vinuela E., Salas M.L.;
 RT "The DNA polymerase-encoding gene of African swine fever virus:
 RL sequence and transcriptional analysis."; Gene 136:103-110(1993).
 RN [17]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93327788; PubMed-8335009;
 RA Simon-Mateo C., Andres G., Vinuela E.;
 RT "Polyprotein processing in African swine fever virus: a novel gene
 RL expression strategy for a DNA virus."; EMBO J. 12:2977-2987(1993).
 RN [18]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93233210; PubMed-8474154;
 RA Prados F.J., Vinuela E., Alcamí A.;
 RT "Sequence and characterization of the major early phosphoprotein p32
 RL of African swine fever virus."; J. Virol. 67:2475-2485(1993).
 RN [19]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-92260660; PubMed-1583732;
 RA Alcamí A., Angulo A., Lopez-Otin C., Munoz M., Freije J.M.,
 RT "Amino acid sequence and structural properties of protein p12, an
 RL African swine fever virus attachment protein."; J. Virol. 66:3860-3868(1992).
 RN [20]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93174976; PubMed-8438592;
 RA Yanez R.J., Vinuela E.;
 RT "African swine fever virus encodes a DNA ligase."; Virology 193:531-536(1993).
 RN [21]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-93174941; PubMed-8382399;
 RA Pena L., Yanez R.J., Revilla Y., Vinuela E., Salas M.L.;
 RT "African swine fever virus guanylyltransferase."; Virology 193:319-328(1993).
 RN [22]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-95159428; PubMed-7856088;

RA Simon-Mateo C., Freije J.M., Andres G., Lopez-Otin C., Vinuela E.;
 RT "Mapping and sequence of the gene encoding protein p17, a major
 RL African swine fever virus structural protein."; Virology 206:1140-1144(1995).
 RN [23]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-92263807; PubMed-1316688;
 RA Garcia-Beato R., Freije J.M., Lopez-Otin C., Blasco R., Vinuela E.,
 RT "A gene homologous to topoisomerase II in African swine fever virus."; Virology 188:938-947(1992).
 RN [24]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BA71V;
 RX MEDLINE-94091056; PubMed-8266720;
 RA Freije J.M., Lain S., Vinuela E., Lopez-Otin C.;
 RT "Nucleotide sequence of a nucleoside triphosphate phosphohydrolase
 Query Match 67.8%; Score 40; DB 12; Length 248;
 Best Local Similarity 63.6%; Pred. No. 20;
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 QY 1 HEAPEPEPIM 11
 Db 227 HEEDEEAEPLI 237
 ID 09VI08 PRELIMINARY; PRT; 1380 AA.
 AC 09VI08;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE CG10132 PROTEIN.
 GN CG10132.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BERKELEY;
 RX MEDLINE-20196006; PubMed-10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotler P.,
 RA Burdus K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Duthin K.J., Evangelista C.C., Ferraz C., Ferrelle S., Fleischmann W.,
 RA Foster C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Idegami C.,
 RA Jaitani M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacleb J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,

RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Strydom R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 Wang Z.-Y., Wasserman D.A., Weinstock G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 DR EMBL: AE003663; AAF53816.1; -
 DR FlyBase: FBgn0032798; CG10132.
 DR InterPro: IPR001680; WD40.
 DR SMART: SM00320; WD40; 2.
 DR PROSITE: PS00678; WD_REPEATS_1; 1.
 DR PROSITE: PS50294; WD_REPEATS_REGION; 1.
 KW Repeat; WD repeat.
 SQ SEQUENCE 1380 AA; 154423 MW; C1928D066450A15B CRC64;

Query Match 67.88; Score 40; DB 5; Length 1380;
 Best Local Similarity 66.78; Pred. No. 1.2e+02;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Oy 1 AEPEAP 9
 Db 179 HDEPTEP 187

RESULT 7
 ID Q9USZ4 PRELIMINARY; PRT; 251 AA.
 AC Q9USZ4;
 DT 01-MAY-2000 (TREMBlrel. 13, Created)
 DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
 DE HYPOTHEICAL 28.0 KDA PROTEIN.
 GN SPB11G11.05.
 OS Schizosaccharomyces pombe (Fission yeast).
 OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;
 OC Schizosaccharomyces.
 OX NCBI_TaxID=4896;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=9728-;
 RA Saunders D., McDougall R.C., Rajandream M.A., Barrell B.G.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AL132717; CAB59807.1; -
 KW Hypothetical protein.
 SQ SEQUENCE 251 AA; 28010 MW; 8018E8325AC65B99 CRC64;

Query Match 66.18; Score 39; DB 3; Length 251;
 Best Local Similarity 87.58; Pred. No. 30;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 2 AEPEAP 9
 Db 185 ESEPEAP 192

RESULT 8
 ID Q9NAB8 PRELIMINARY; PRT; 304 AA.
 AC Q9NAB8;
 DT 01-OCT-2000 (TREMBlrel. 15, Created)
 DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
 DE Y33FAB.16 PROTEIN.
 GN Y33FAB.16;
 OS *Caenorhabditis elegans*.

OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
 OC Rhabditidae; Pelodierinae; *Caenorhabditis*.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Smye R.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=99069613; PubMed=9851916;
 RA none;
 RT "Genome sequence of the nematode *C. elegans*: A platform for
 RT investigating biology.";
 RL Science 282:2012-2018(1998).
 DR EMBL: AL132949; CAB61086.1; -
 SQ SEQUENCE 304 AA; 34146 MW; 20AB91D8BB137A76 CRC64;

Query Match 66.18; Score 39; DB 5; Length 304;
 Best Local Similarity 87.58; Pred. No. 37;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 3 AEPEAP 10
 Db 80 AEPEAP 87

RESULT 9
 ID Q9NAA2 PRELIMINARY; PRT; 340 AA.
 AC Q9NAA2;
 DT 01-OCT-2000 (TREMBlrel. 15, Created)
 DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
 DT 01-MAR-2001 (TREMBlrel. 16, Last annotation update)
 DE Y33FAB.17 PROTEIN.
 GN Y33FAB.17.
 OS *Caenorhabditis elegans*.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
 OC Rhabditidae; Pelodierinae; *Caenorhabditis*.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Smye R.;
 RL Submitted (OCT-1999) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=99069613; PubMed=9851916;
 RA none;
 RT "Genome sequence of the nematode *C. elegans*: A platform for
 RT investigating biology.";
 RL Science 282:2012-2018(1998).
 DR EMBL: AL132949; CAB70107.1; -
 SQ SEQUENCE 340 AA; 36879 MW; 2EC01E30582C9E3A CRC64;

Query Match 66.18; Score 39; DB 5; Length 340;
 Best Local Similarity 87.58; Pred. No. 42;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 3 AEPEAP 10
 Db 247 AEPEAP 254

RESULT 10
 ID Q96979 PRELIMINARY; PRT; 408 AA.
 AC Q96979;
 DT 01-MAY-1999 (TREMBlrel. 10, Created)
 DT 01-MAY-1999 (TREMBlrel. 10, Last sequence update)
 DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
 DE TRANSLATION ELONGATION FACTOR 1-ALPHA (FRAGMENT).
 GN TEF1.

OS Paranophrys carnivora.
 OC Eukaryota; Alveolata; Ciliophora; Oligohymenophorea; Scuticociliatida;
 OC Paranophryidae; Paranophrys.
 NX NCBI_TaxID=85900;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=99152616; PubMed=10028290;
 RA Morella D., Le Guyader H., Philippe H.;
 RT "Unusually high evolutionary rate of the elongation factor 1 alpha
 RT genes from the Ciliophora and its impact on the phylogeny of
 RT eukaryotes.";
 RL Mol. Biol. Evol. 16:234-245(1999).
 CC -I- SIMILARITY: TO GTP-BINDING ELONGATION FACTOR FAMILY.
 DR EMBL: AF056103; AAD03258.1; -
 DR HSSP: Q01698; 1PTT.
 DR InterPro: IPR000795; GTP_EFTU.
 DR Pfam: PF00009; GTP_EFTU; 1.
 DR PRINTS: PR00315; ELONGATNCT.
 DR PROSITE: PS00301; EFACOR_GTP; 1.
 KW Elongation factor; GTP-binding; Protein biosynthesis.
 FT NON_TER 1
 FT NON_TER 408
 SQ SEQUENCE 408 AA; 4506 MW; 384973BB7F3F5FAL CRC64;

Query Match 66.1%; Score 39; DB 5; Length 408;
 Best Local Similarity 77.8%; Pred. No. 50;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 HEAPEAP 9
 Db 269 HESLEAP 277

RESULT 11
 ID O9MAY3 PRELIMINARY; PRT; 417 AA.
 AC O9MAY3;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE CG12497 PROTEIN.
 GN EG:BACR25B3.2 OR CG12497.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 NX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H.C., Blaise R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Mills G.L.G.,
 RA Abrell J.F., Adiyanti A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferriera S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glasser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostali D., Houston K.A., Howland J.J., Wei M.-H., Ibeagwa C.,
 RA Jostali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,

RA Lasko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstock G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster.";
 RL Science 287:2185-2195(2000).
 CC -I- SIMILARITY: TO LOW DENSITY LIPOPROTEIN (LDL) RECEPTOR CLASS A
 CC (LDLRA) DOMAIN.
 DR EMBL: AE003424; AAF45787.1; -
 DR HSSP: Q07954; 1CR8.
 DR FlyBase: FBgn0040379; EG:BACR25B3.2.
 DR InterPro: IPR002172; LDL_recept_A.
 DR Pfam: PF00057; LDL_recept_a; 1.
 DR PRINTS: PR00261; LDLRECEPTOR.
 DR SMART: SM00192; LDLA; 2.
 DR PROSITE: PS01209; LDLRA_1; 1.
 DR PROSITE: PS50068; LDLRA_2; 2.
 KW Glycoprotein.
 SQ SEQUENCE 417 AA; 48624 MW; 2F1CA7F440D1DD01 CRC64;

Query Match 66.1%; Score 39; DB 5; Length 417;
 Best Local Similarity 87.5%; Pred. No. 51;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
 Db 382 ESEPEAP 389

RESULT 12
 ID O9NEF8 PRELIMINARY; PRT; 435 AA.
 AC O9NEF8;
 DT 01-OCT-2000 (TREMBLrel. 15, Created)
 DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE EG:BACR25B3.2 OR CG12497.
 GN EG:BACR25B3.2 OR CG12497.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 NX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Murphy L., Harris D., Barrell B.;
 RT "Sequencing the distal X chromosome of Drosophila melanogaster.";
 RT Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Benos P.;
 RT Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
 CC -I- SIMILARITY: TO LOW DENSITY LIPOPROTEIN (LDL) RECEPTOR CLASS A
 CC (LDLRA) DOMAIN.
 DR EMBL: AL18972; CAB72287.1; -
 DR FlyBase: FBgn0040379; EG:BACR25B3.2.
 DR InterPro: IPR002172; LDL_recept_A.
 DR Pfam: PF00057; LDL_recept_a; 1.
 DR PRINTS: PR00261; LDLRECEPTOR.
 DR SMART: SM00192; LDLA; 2.
 DR PROSITE: PS01209; LDLRA_1; 1.

TRANS-ACTING TRANSCRIPTIONAL PROTEIN ICP4
DE (IMMEDIATE-EARLY PROTEIN IE62).
GN 62.
OS Cercopithecine herpesvirus 9 (simian varicella virus).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicelloviruses.
OX NCBI_TaxID=35246;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=94023599; PubMed=8212583;
RA Clarke P., Brunschwig A., Gildea D.H.;
RT "DNA sequence of a simian varicella virus gene that encodes a
RT homologue of varicella zoster virus IE62 and herpes simplex virus
RT ICP4."
RL Virology 197:45-52(1993).
CC -1- FUNCTION: THIS IE PROTEIN IS A MULTIFUNCTIONAL PROTEIN CAPABLE OF
CC MIGRATING TO THE NUCLEUS, BINDING TO DNA, TRANS-ACTIVATING OTHER
CC VIRAL GENES, AND AUTOREGULATING ITS OWN SYNTHESIS.
CC -1- SUBCELLULAR LOCATION: NUCLEUS OF INFECTED CELLS.
CC -1- PTM: A LONG STRETCH OF SERINE RESIDUES MAY BE A MAJOR SITE OF
CC PHOSPHORYLATION
CC -1- SIMILARITY: BELONGS TO THE ICP4/IE140/IE180 FAMILY OF PROTEINS.
CC EMBL: L20759; AAA03549.1; -.
DR InterPro: IPR000923; Cooper_blue1.
DR PROSITE: PS00196; CQPPER_BLUE; UNKNOWN_1.
KW Early protein; Transcription regulation; Trans-acting factor;
KW DNA-binding; Nuclear protein.
SQ SEQUENCE 1277 AA; 136978 MW; BB92AA8C4DCBD9D CRC64;

Query Match 66.18; Score 39; DB 12; Length 1277;
Best local similarity 70.0%; Pred. No. 1.6e+02;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAEPEAPEIM 11
|:|||||:
Db 1106 EDPPEAPEPII 1115

Search completed: January 7, 2002, 08:47:23
Job time: 411 sec

GenCore version 4.5
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OM protein - protein search, using sw.model

Run on: January 4, 2002, 08:41:02 ; Search time 18.1 seconds
(without alignments)
22.282 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAEPFAPRIM 11

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 100059 seqs, 36664827 residues

Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: SwissProt_39

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	42	71.2	144	1 AP22_APIME	P35581 apis mellif
2	42	71.2	283	1 AP73_APIME	Q06602 apis mellif
3	40	67.8	229	1 CGB1_CRIO	Q08301 cricetus
4	39	66.1	486	1 HSI_HUMAN	P13177 homo sapien
5	38	64.4	25	1 NP4_HUMAN	P18078 homo sapien
6	38	64.4	168	1 AP14_APIME	Q06601 apis mellif
7	38	64.4	248	1 GRL1_RAT	Q06605 ratus norv
8	38	64.4	256	1 PRN3_HUMAN	P24158 homo sapien
9	38	64.4	332	1 H630_YEAST	P25619 saccharomyc
10	37	62.7	226	1 DDN1_BOVIN	P80219 bos taurus
11	37	62.7	245	1 MCT1_SHEEP	P80931 ovis aries
12	37	62.7	246	1 GRAB_HUMAN	P20718 homo sapien
13	37	62.7	247	1 TONB_HUMAN	P10144 h girazyme
14	37	62.7	247	1 TONB_SERNA	P26185 serralia ma
15	37	62.7	248	1 NKPI_RAT	P18291 ratus norv
16	37	62.7	251	1 MCT3_SHEEP	O46683 ovis aries
17	37	62.7	269	1 YAS3_ARCFU	O29209 archaeoglob
18	37	62.7	376	1 XYNA_BACOV	P49942 bacteroides
19	37	62.7	483	1 PHR_STINE	P05327 synechococ
20	37	62.7	601	1 PAGT_CAEEL	P36678 caenorhabdi
21	36	61.0	182	1 YSM2_CAEEL	Q10122 caenorhabdi
22	36	61.0	239	1 TONB_ECOLI	P02929 escherichia
23	36	61.0	242	1 TONB_SALTY	P25945 salmonella
24	36	61.0	255	1 TONB_YEREN	O05740 yersinia en
25	36	61.0	277	1 PS12_ARATH	O23712 arabidopsis
26	36	61.0	372	1 NK11_HUMAN	Q15759 homo sapien
27	36	61.0	423	1 CGB1_RAT	P30277 ratus norv
28	36	61.0	565	1 MOT8_MOUSE	O70324 mus musculu
29	36	61.0	1195	1 KDGD_HUMAN	Q16760 homo sapien
30	35	59.3	171	1 ZUR_ECOLI	P32692 escherichia
31	35	59.3	292	1 SUB1_YEAST	P54000 saccharomyc
32	35	59.3	330	1 XYN2_BACST	P45703 bacillus st
33	35	59.3	433	1 CGB1_HUMAN	P14635 homo sapien

34	35	59.3	465	1 G3BP_MOUSE	P97855 mus musculu
35	35	59.3	466	1 G3BP_HUMAN	Q13283 homo sapien
36	35	59.3	474	1 SY65_DROME	P21521 drosophila
37	35	59.3	486	1 HSL_MOUSE	P49710 mus musculu
38	35	59.3	613	1 MOT8_HUMAN	P36021 homo sapien
39	35	59.3	754	1 RAD4_YEAST	P14736 saccharomyc
40	35	59.3	788	1 TRS1_HCVMA	P09695 human cytom
41	35	59.3	2749	1 IP3R_MOUSE	P11881 mus musculu
42	35	59.3	2749	1 IP3R_RAT	P29994 ratus norv
43	34	57.6	202	1 B3G1_MOUSE	O96w73 m galactosy
44	34	57.6	302	1 RS3_HALLA	P15009 halobacteri
45	34	57.6	334	1 B3G1_RAT	O35789 r galactosy

ALIGNMENTS

RESULT	ID	AP22_APIME	STANDARD	PRT	144 AA
AC	P35581	P11525; P11526;			
DT	01-OCT-1989	(Rel. 12, Created)			
DT	01-JUN-1994	(Rel. 29, Last sequence update)			
DT	01-JUN-1994	(Rel. 29, Last annotation update)			
DE	APIDAECIN PRECURSOR, TYPE 22.				
OS	Apis mellifera (Honeybee).				
OC	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;				
OC	Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata;				
OX	NCBI_TaxID=7460;				
RM	11				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=93223697; PubMed=8467807;				
RA	Casteels-Josson K., Capaci T., Casteels P., Tempst P.;				
RT	"Apidaecin multiprotein precursor structure: a putative mechanism for				
RT	amplification of the insect antibacterial response.";				
RL	EMBO J. 12:1569-1578(1993).				
RM	12				
RP	SEQUENCE (APIDAECIN IA/IB).				
RC	TISUP-Hemolymph;				
RX	MEDLINE=90005446; PubMed=2676519;				
RA	Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;				
RT	"Apidaecins: antibacterial peptides from honeybees.";				
RL	EMBO J. 8:2387-2391(1989).				
CC	- FUNCTION: APIDAECIN HAVE BACTERICIDAL ACTIVITY; PREDOMINANTLY				
CC	AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL				
CC	PROLIFERATION.				
CC	-----				
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration				
CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -				
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CC	modified and this statement is not removed. Usage by and for commercial				
CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/				
CC	or send an email to license@sib-sib.ch).				
CC	-----				
DR	EMBL; X72576; CAAS1168.1; -				
DR	PIR; S05383; S05383.				
DR	PIR; S06675; S06675.				
DR	PIR; S35331; S35331.				
DR	InterPro: IPR001979; Apidaecin.				
DR	Pfam: PF00807; Apidaecin; 4.				
KW	Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;				
KW	Cleavage on pair of basic residues; Repeat.				
FT	SIGNAL	1	19	POTENTIAL.	
FT	PROPEP	35	42		
FT	PEPTIDE	43	60	APIDAECIN IB.	
FT	PROPEP	63	70		
FT	PEPTIDE	71	88	APIDAECIN IB.	
FT	PROPEP	91	98		
FT	PEPTIDE	99	116	APIDAECIN IB.	
FT	PROPEP	119	126		
FT	PEPTIDE	127	144	APIDAECIN IA.	

SQ SEQUENCE 144 AA; 16539 MW; 6FA1AD74CB71108D CRC64;

Query Match
Best Local Similarity 71.2%; Score 42; DB 1; Length 144;
Matches 8; Conservative 100.0%; Pred. No. 1.4;
Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
|111111|
DB 35 EAEPEAP 42

RESULT 2
ID AP73.APIME STANDARD; PRT: 283 AA.
AC 006602: P11525; P11526;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-JUN-1994 (Rel. 29, Last annotation update)
DE APIDAECIN PRECURSOR, TYPE 73 (FRAGMENT).
GN APID73.
OS Apis mellifera (Honeybee).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata;
OC Apoidea; Apidae; Apis.
OX NCBI_TaxID=7460;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=93223697; PubMed=8467807;
RA Casteels-Josson K., Capaci T., Casteels P., Tempst P.;
RT "Apidaecin multipetide precursor structure: a putative mechanism for
amplification of the insect antibacterial response.";
RL EMO J. 8:2387-2391(1989).
RN [2]
RP SEQUENCE OF APIDAECIN IA/IB.
RC TISSUE=Hemolymph;
RX MEDLINE=90005446; PubMed=2676519;
RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
RT "Apidaecins: antibacterial peptides from honeybees.";
RL EMO J. 8:2387-2391(1989).
CC -I- FUNCTION: APIDAECIN HAVE BACTERICIDAL ACTIVITY: PREDOMINANTLY
AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
PROLIFERATION.
CC -----
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or send an email to license@sib-sib.ch).
CC -----
CC EMBL: X72577; CAA51169.1; -
DR PIR: S05383; S05383.
DR PIR: S06675; S06675.
DR PIR: S35332; S35332.
DR InterPro: IPR001979; Apidaecin.
DR Pfam: PF00807; Apidaecin.9.
DR ProDom: PD153432; Apidaecin.2.
KW Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
KW Cleavage on pair of basic residues; Repeat.
FT SIGNAL 1 18 POTENTIAL.
FT PROPEP 34 41
FT PROPEP 42 59 APIDAECIN IB.
FT PROPEP 62 69
FT PROPEP 70 87 *APIDAECIN IB.
FT PROPEP 90 97
FT PROPEP 98 115 APIDAECIN.
FT PROPEP 118 135
FT PROPEP 126 143 APIDAECIN IB.
FT PROPEP 146 153
FT PROPEP 154 171 APIDAECIN.
FT *PEPTIDE

FT PROPEP 174 182
FT PEPTIDE 183 199 APIDAECIN IB.
FT PROPEP 202 209
FT PEPTIDE 210 227 APIDAECIN IB.
FT PROPEP 230 237
FT PEPTIDE 238 255 APIDAECIN IB.
FT PROPEP 258 265
FT PEPTIDE 266 283
SQ SEQUENCE 283 AA; 32695 MW; 4EA5FEDEC05E142B CRC64;

Query Match
Best Local Similarity 71.2%; Score 42; DB 1; Length 283;
Matches 8; Conservative 100.0%; Pred. No. 2.9;
Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
|111111|
DB 62 EAEPEAP 69

RESULT 3
ID CGBL_CRILLO STANDARD; PRT: 429 AA.
AC 008301;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-OCT-1994 (Rel. 30, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE G2/MITOTIC-SPECIFIC CYCLIN B1.
GN CCNB1.
OS Cricetus longicaudatus (Long-tailed hamster) (Chinese hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;
OC Cricetulus.
OX NCBI_TaxID=10030;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=ovary;
RX MEDLINE=94095439; PubMed=8270434;
RA Markiewicz D.A., McKenna W.G., Flick M.B., Maity A., Muschel R.J.;
RT "The effects of radiation on the expression of a newly cloned and
characterized rat cyclin B mRNA.";
RL Int. J. Radiat. Oncol. Biol. Phys. 28:135-144(1994).
RN [2]
RP -I- FUNCTION: ESSENTIAL FOR THE CONTROL OF THE CELL CYCLE AT THE G2/M
(MITOSIS) TRANSITION.
CC -I- SUBUNIT: INTERACTS WITH THE CDC2 PROTEIN KINASE TO FORM A
SERINE/THREONINE KINASE HOLOENZYME COMPLEX ALSO KNOWN AS
MATURATION PROMOTING FACTOR (MPF). THE CYCLIN SUBUNIT IMPARTS
SUBSTRATE SPECIFICITY TO THE COMPLEX.
CC -I- DEVELOPMENTAL STAGE: ACCUMULATES STEADILY DURING G2 AND IS
ABRUPTLY DESTROYED AT MITOSIS.
CC -I- SIMILARITY: BELONGS TO THE CYCLIN FAMILY. CYCLIN AB SUBFAMILY.
CC -----
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CC -----
CC EMBL: X64588; CAA5876.1; -
DR PIR: S34224; S34224.
DR HSP: P20248; IJST.
DR InterPro: IPR000553; Cyclin.
DR Pfam: PF00134; cyclin.1.
DR SMART: SM00385; CYCLIN; 2.
DR PROSITE: PS00292; CYCLIN; 1.
KW Cyclin; Cell cycle; Cell division; Mitosis.
KW DOMAIN 51 86 LYS-RICH.
SQ SEQUENCE 429 AA; 48062 MW; 6E0BAE7511A678B7 CRC64;

Query Match 67.8%; Score 40; DB 1; Length 429;

Best Local Similarity 70.0%; Pred. No. 9.6;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Oy 2 HEAPEPEPIM 11
| | | | | : |
Db 98 EPEPEPEPVM 107

RESULT 4

HSL_HUMAN STANDARD: PRT; 486 AA.
ID HSL_HUMAN
AC P14317;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE HEMATOPOIETIC LINEAGE CELL SPECIFIC PROTEIN (HEMATOPOIETIC CELL-
SPECIFIC LYN SUBSTRATE 1) (LCKBPL1).
GN HCLSI OR HSL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90067934; PubMed=2587259;
RA Kitamura D., Kaneko H., Miyagoe Y., Ariyasu T., Watanabe T.;
RT "Isolation and characterization of a novel human gene expressed
specifically in the cells of hematopoietic lineage.";
RL Nucleic Acids Res. 17:9367-9379(1989).
RN [2]
RP SEQUENCE OF 97-108; 193-201 AND 240-248.
RX MEDLINE=9631348; PubMed=8713105;
RA Egerton M., Moritz R.L., Druker B., Kelso A., Simpson R.J.;
RT "Identification of the 70KD heat shock cognate protein (Hsc70) and
alpha-actinin-1 as novel phosphotyrosine-containing proteins in T
lymphocytes.";
RL Biochem. Biophys. Res. Commun. 224:666-674(1996).
RN [3]
RP BINDING TO HAX-1 PROTEIN.
RX MEDLINE=97211841; PubMed=9059808;
RA Suzuki Y., Demoliere C., Kitamura D., Takeshita H., Deuschle U.,
Watanabe T.;
RT "HAX-1, a novel intracellular protein, localized on mitochondria,
directly associates with HSL, a substrate of src family tyrosine
kinases.";
RL J. Immunol. 158:2736-2744(1997).
RN [4]
RP FUNCTION: SUBSTRATE OF THE ANTIGEN RECEPTOR-COUPLED TYROSINE
KINASE. PLAYS A ROLE IN ANTIGEN RECEPTOR SIGNALING FOR BOTH
CLONAL EXPANSION AND DELETION IN LYMPHOID CELLS. DIRECTLY
ASSOCIATES WITH HAX-1, THROUGH BINDING TO ITS C-TERMINAL REGION.
MAY ALSO BE INVOLVED IN THE REGULATION OF GENE EXPRESSION.
CC -1 SUBUNIT: ASSOCIATES WITH THE SH2 AND SH3 DOMAINS OF LCK.
CC -1 SUBCELLULAR LOCATION: MITOCHONDRIAL (PROBABLE).
CC -1 TISSUE SPECIFICITY: EXPRESSED ONLY IN TISSUES AND CELLS OF
HEMATOPOIETIC ORIGIN.
CC -1 DEVELOPMENTAL STAGE: EXPRESSED IN EARLY STAGE OF MYELOID AND
ERYTHROID DIFFERENTIATION.
CC -1 PM: PHOSPHORYLATED BY LYN RAPIDLY AFTER CROSSLINKING OF SURFACE
TCR ON B CELLS.
CC -1 SIMILARITY: TO CHICKEN P80/85 PROTEINS (CONTACTIN).
CC -1 SIMILARITY: CONTAINS 1 SH3 DOMAIN.
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CC -----
DR EMBL: X16663; CA34651.1; -;
DR PIR: S07633; S07633.
DR HSSP: P29355; 1SEM.

DR MM: 601306; -;
DR InterPro: IPR003134; HSL_rep.
DR InterPro: IPR001452; SH3
DR Pfam: PF02218; HSL_rep; 4.
DR Pfam: PF00018; SH3; 1.
DR PRINTS: PR00452; SH3DOMAIN.
DR SMART: SM00326; SH3; 1.
DR PROSITE: PS50002; SH3; 1.
KW Repeat: SH3 domain; Phosphorylation.
FT DOMAIN 27 66 INVOLVED IN HAX-1 BINDING.
FT DOMAIN 81 214 3.5 X 37 AA TANDEM REPEATS.
FT REPEAT 81 116 1.
FT REPEAT 117 153 2.
FT REPEAT 154 190 3.
FT REPEAT 191 214 4 (INCOMPLETE).
FT DOMAIN 428 486 SH3.
FT CONFLICT 241 242 KF -> PK (IN REF. 2).
SQ SEQUENCE 486 AA; 53998 MW; 61AE637157DF5DF2 CRC64;

Query Match 66.1%; Score 39; DB 1; Length 486;
Best Local Similarity 77.8%; Pred. No. 16;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Oy 1 HEAPEPEP 9
| | | | | : |
Db 360 YEAPPEPEP 368

RESULT 5

NP4_HUMAN STANDARD: PRT; 25 AA.
ID NP4_HUMAN
AC P18078;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE NEUTROPHIL PROTEINASE 4 (EC 3.4.21.-) (NP-4) (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE.
RX MEDLINE=91025622; PubMed=2121162;
RA Ohlsson K., Linder C., Rosengren M.;
RT "Monoclonal antibodies specific for neutrophil proteinase 4.
Production and use for isolation of the enzyme.";
RL Biol. Chem. Hoppe-Seyler 371:549-555(1990).
CC -1 TISSUE SPECIFICITY: NEUTROPHILS.
CC -1 SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
TRYPSIN FAMILY.
CC HSSP: P24158; 1FUJ.
DR MEROPS: S01.134; -;
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; Trypsin; 1.
DR PROSITE: PS50240; TRYPSIN_DOM; PARTIAL.
DR PROSITE: PS00134; TRYPSIN_HIS; PARTIAL.
DR PROSITE: PS00135; TRYPSIN_SER; PARTIAL.
KW Hydrolyase; Serine protease; Glycoprotein.
FT NON_TER 25
SQ SEQUENCE 25 AA; 2606 MW; B1CB2038274575B0 CRC64;

Query Match 64.4%; Score 38; DB 1; Length 25;
Best Local Similarity 54.5%; Pred. No. 1.2;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Oy 1 HEAPEPEPIM 11
| | | | | : |
Db 5 HEADPHSRPYM 15

RESULT 6

AP14_APIME STANDARD: PRT: 168 AA.
 ID AP14_APIME STANDARD: PRT: 168 AA.
 AC 006601; P11525; P11526; P11527;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 01-JUN-1994 (Rel. 29, Last annotation update)
 DE APIDAEIN PRECURSOR, TYPE 14.
 GN APID14.
 OS Apis mellifera (Honeybee).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pelegroneta; Neoptera; Endopterygota; Hymenoptera; Apoecrita; Aculeata;
 OC Apoidea; Apidae; Apis.
 OX NCBI_TaxID=7460;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93223697; PubMed=8467807;
 RA Casteels J., Jossion K., Capact T., Casteels P., Tempst P.;
 RT "Apidaecin multipeptide precursor structure: a putative mechanism for
 amplification of the insect antibacterial response.";
 RL EMBO J. 12:1569-1578(1993).
 RN [2]
 RP SEQUENCE OF APIDAEIN IN/IB/II.
 RC TISSUE=Hemolymph;
 RX MEDLINE=90005446; PubMed=2676519;
 RA Casteels P., Ampe C., Jacobs F., Vaeck M., Tempst P.;
 RT "Apidaecins: antibacterial peptides from honeybees.";
 RL EMBO J. 8:2387-2391(1989).
 CC -1- FUNCTION: APIDAEIN HAS BACTERICIDAL ACTIVITY. PREDOMINANTLY
 AGAINST GRAM-NEGATIVE BACTERIA. THEY SEEM TO INTERFERE WITH CELL
 PROPAGATION.
 CC -----
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 CC -----
 DR EMBL: X72575; CAA51167.1; -
 DR PIR: S05383; S05383.
 DR PIR: S06675; S06675.
 DR PIR: S06676; S06676.
 DR PIR: S35330; S35330.
 DR InterPro: IPR001979; Apidaecin.
 DR Pfam: PF00807; Apidaecin; 5.
 DR ProDom: PD153432; Apidaecin; 1.
 KM Insect immunity; Antibiotic; Hemolymph; Signal; Multigene family;
 KM Cleavage on pair of basic residues; Repeat.
 KM SIGNAL 1
 FT 19
 FT PROPEP 35 42
 FT ACT_SITE 60
 FT ACT_SITE 63 70
 FT ACT_SITE 71 88
 FT PROPEP 91 98
 FT PEPTIDE 99 116
 FT PROPEP 119 124
 FT PEPTIDE 125 142
 FT PROPEP 145 150
 FT PEPTIDE 151 168
 FT SEQUENCE 168 AA; 19380 MW; 594B931254C04A37 CRC64;
 SO APIDAEIN IA.
 Query Match 64.4%; Score 38; DB 1; Length 168;
 Best Local Similarity 87.5%; Pred. No. 8.3;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

GR1L_RAT STANDARD: PRT: 248 AA.
 ID GR1L_RAT STANDARD: PRT: 248 AA.
 AC 006605;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE GRANZYME-LIKE PROTEIN I PRECURSOR (EC 3.4.21.-).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93285341; PubMed=8508925;
 RA Amerik A.Y., Jarovoi S.V., Grigorenko V.G., Antonov V.K.;
 RT "Identification, sequence analysis, and characterization of cDNA
 clones encoding two granzyme-like serine proteinases from rat
 duodenum.";
 RL FEBS Lett. 324:226-230(1993).
 CC -1- FUNCTION: THIS ENZYME IS NECESSARY FOR TARGET CELL LYSIS IN
 CELL-MEDIATED IMMUNE RESPONSES.
 CC -1- TISSUE SPECIFICITY: DUODENUM.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY. STRONGEST TO GRANZYMES AND TO MAST CELL PROTEASES.
 CC -----
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 CC -----
 DR EMBL: X66693; CAA47235.1; -
 DR HSSP: P04187; 2CPL.
 DR MEROPS: S01.136; -
 DR InterPro: IPR001314; Chymotrypsin.
 DR InterPro: IPR001254; Trypsin.
 DR Pfam: PF00089; trypsin; 1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR SMART: SM00020; TRYP-Spec; 1.
 DR PROSITE: PS50240; TRYPSIN_DOM; 1.
 DR PROSITE: PS00134; TRYPSIN_HIS; 1.
 DR PROSITE: PS00135; TRYPSIN_SER; 1.
 KM Hydrolyase; Serine protease; Zymogen; Signal; Glycoprotein.
 FT SIGNAL 1
 FT PROPEP 19 20
 FT CHAIN 21 248
 FT ACT_SITE 65 65
 FT ACT_SITE 109 109
 FT ACT_SITE 204 204
 FT DISULFID 50 66
 FT DISULFID 143 210
 FT DISULFID 174 189
 FT CARBOHYD 72 72
 FT SEQUENCE 248 AA; 27235 MW; 036C81B43A8B972 CRC64;
 SO N-LINKED (GLCNAC...) (POTENTIAL).
 Query Match 64.4%; Score 38; DB 1; Length 248;
 Best Local Similarity 54.5%; Pred. No. 12;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 2 EAPPEAP 9
 DB 35 EAKPEAP 42
 RESULT 7
 DT 01-APR-1990 (Rel. 14, Created)

PRN3_HUMAN STANDARD: PRT: 256 AA.
 ID PRN3_HUMAN STANDARD: PRT: 256 AA.
 AC P24158; P15637;
 DT 01-APR-1990 (Rel. 14, Created)

DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE MYELOBLASTIN PRECURSOR (PC 3.4-21.76) (LEUKOCYTE PROTEINASE 3) (PR-3)
 GN (PR3) (GSP7) (WEGENER'S AUTOANTIGEN) (P29) (C-ANCA ANTIGEN).
 CN PRTN3 OR MBN.
 OS Homo sapiens (human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 NC NCBL_Taxid=9606;
 [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92021028; PubMed=1681549;
 RA Labbaye C., Musette P., Cayre Y.E.;
 RT "Wegener autoantigen and myeloblastin are encoded by a single mRNA.";
 RL Proc. Natl. Acad. Sci. U.S.A. 88:9253-9256(1991).
 [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91079774; PubMed=2258701;
 RA Campenelli D., Melchior M., Fu Y., Nakata M., Shuman H., Nathan C.,
 Gabay J.E.;
 RT "Cloning of cDNA for proteinase 3: a serine protease, antibiotic, and
 RT autoantigen from human neutrophils.";
 RL J. Exp. Med. 172:1709-1715(1990).
 [4]
 RP SEQUENCE OF 1-20 AND 22-256 FROM N.A.
 RX MEDLINE=92390417; PubMed=1518849;
 RA Zimmer M., Medcalf R.L., Fink T.M., Maltmann C., Lichter P.,
 Jenne D.E.;
 RT "Three human elastase-like genes coordinately expressed in the
 RT myelomonocyte lineage are organized as a single genetic locus on
 RT 19pter.";
 RL Proc. Natl. Acad. Sci. U.S.A. 89:8215-8219(1992).
 [5]
 RP SEQUENCE OF 42-256 FROM N.A.
 RX MEDLINE=90090622; PubMed=2596267;
 RA Borjes D., Raynal M.-C., Solomon D.H., Darzynkiewicz Z., Cayre Y.E.;
 RT "Down-regulation of a serine protease, myeloblastin, causes growth
 RT arrest and differentiation of promyelocytic leukemia cells.";
 RL Cell 59:959-968(1989).
 [6]
 RP SEQUENCE OF 28-67 AND 228-244.
 RX MEDLINE=91236723; PubMed=2033050;
 RA Rao N.V., Wehner N.G., Marshall B.C., Gray W.R., Gray B.H.,
 Hoidal J.R.;
 RT "Characterization of proteinase-3 (PR-3), a neutrophil serine
 RT proteinase. Structural and functional properties.";
 RL J. Biol. Chem. 266:9540-9548(1991).
 [7]
 RP SEQUENCE OF 28-47.
 RX MEDLINE=89315847; PubMed=2501794;
 RA Gabay J.E., Scott R.W., Campenelli D., Griffith J., Wilde C.,
 Maria M.N., Seeger M., Nathan C.F.;
 RT "Antibiotic proteins of human polymorphonuclear leukocytes.";
 RL Proc. Natl. Acad. Sci. U.S.A. 86:5610-5614(1989).
 [8]
 RP SEQUENCE OF 28-47 AND 196-219.
 RX MEDLINE=90130450; PubMed=2404977;
 RA Wilde C.G., Snable J.L., Griffith J.E., Scott R.W.;
 RT "Characterization of two azurophilic granule proteases with active-site
 RT homology to neutrophil elastase.";
 RL J. Biol. Chem. 265:2038-2041(1990).
 [9]

RP SEQUENCE OF 28-48, AND IDENTITY OF WEGENER'S AUTOANTIGEN WITH PR-3.
 RX MEDLINE=9032035; PubMed=2377228;
 RA Jenne D.E., Tschopp J., Luedemann J., Utecht B., Gross W.L.;
 RT "Wegener's autoantigen decoded.";
 RL Nature 346:520-520(1990).
 [10]
 RP IDENTITY OF WEGENER'S AUTOANTIGEN WITH PROTEINASE 3.
 RX MEDLINE=91055123; PubMed=2242436;
 RA Gupta S.K., Niles J.L., McCluskey R.T., Annaut M.A.;
 RT "Identity of Wegener's autoantigen (p29) with proteinase 3 and
 RT myeloblastin.";
 RL Blood 76:2162-2162(1990).
 [11]
 RP X-RAY CRYSTALLOGRAPHY (2.2 ANGSTROMS).
 RX MEDLINE=96346178; PubMed=8757293;
 RA Fujinaga M., Charnala M.M., Halenbeck R., Kothe K., James M.N.G.;
 RT "The crystal structure of PR3, a neutrophil serine proteinase antigen
 RT of Wegener's granulomatosis antibodies.";
 RL J. Mol. Biol. 261:267-278(1996).
 CC -1- FUNCTION: POLYMORPHONUCLEAR LEUKOCYTE SERINE PROTEASE THAT
 CC DEGRADES ELASTIN, FIBRONECTIN, LAMININ, VITRONECTIN, AND COLLAGEN
 CC TYPES I, III, AND IV (IN VITRO) AND CAUSES EMPHYSEMA WHEN
 CC ADMINISTERED BY TRACHEAL INSUFFLATION TO HAMSTERS.
 CC -1- CATALYTIC ACTIVITY: HYDROLYSIS OF PROTEINS, INCLUDING ELASTIN, BY
 CC PREFERENTIAL CLEAVAGE: ALA-I-XAA > VAL-I-XAA.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 CC TRYPSIN FAMILY. ELASTASE SUBFAMILY.
 CC -----
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 CC -----
 DR EMBL: X56132; CAA39598.1; -;
 DR EMBL: AC004799; AAC18958.1; -;
 DR EMBL: M75154; AAA59558.1; -;
 DR EMBL: M96839; AAB59493.1; -;
 DR EMBL: M96838; AAB59493.1; JOINED.
 DR EMBL: M96837; AAB59493.1; JOINED.
 DR EMBL: X55668; CAA39203.1; -;
 DR EMBL: M29142; AAA36342.1; -;
 DR EMBL: M96628; AAB59364.1; -;
 DR PIR: A43983; PRH03.
 DR PDB: 1F0J; 11-JUL-96.
 DR MEROPS: S01.134; -;
 DR MIM: 177020; -;
 DR InterPro: IPR001254; Chymotrypsin.
 DR Pfam: PF00089; Trypsin.1.
 DR PRINTS: PR00722; CHYMOTRYPSIN.
 DR SMART: SM0020; TRYP-SPC.1.
 DR PROSITE: PS50240; TRYPSIN_DOM.1.
 DR PROSITE: PS00134; TRYPSIN_HIS.1.
 DR PROSITE: PS00135; TRYPSIN_SER.1.
 KW Hydrolyase; Serine protease; Glycoprotein; Zymogen; Signal;
 KW 3D-structure.
 FT SIGNAL 1 25
 FT PROPEP 26 27
 FT CHAIN 28 248
 FT PROPEP 249 256
 FT ACT_SITE 71 71
 FT ACT_SITE 118 118
 FT ACT_SITE 203 203
 FT CARBOHYD 129 129
 FT CARBOHYD 174 174
 FT DISULFID 56 72
 FT DISULFID 152 209
 FT DISULFID 182 188
 FT DISULFID 199 224
 FT CONFLICT 2 2
 A -> R (IN REF. 3).

FT CONFLICT 46 46 Q -> E (IN REF. 7 AND 8).
 FT CONFLICT 64 64 S -> D (IN REF. 6).
 FT CONFLICT 70 70 A -> P (IN REF. 1).
 FT CONFLICT 119 119 V -> I (IN REF. 1 AND 5).
 FT CONFLICT 135 136 AT -> TS (IN REF. 1 AND 5).
 FT CONFLICT 255 255 MISSING (IN REF. 3).
 SQ SEQUENCE 256 AA; 27807 MW; CBECA36D8C4B2A40 CRC64;

Query Match 64.4%; Score 38; DB 1; Length 256;
 Best Local Similarity 54.5%; Pred. No. 13;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIIM 11
 DB 32 HEAPHSPRPM 42

RESULT 9
 HS30_YEAST STANDARD; PRT; 332 AA.
 AC P25619; 004556;
 DT 01-MAY-1992 (Rel. 22, Created)
 DT 01-MAY-1992 (Rel. 22, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE 30 KDA HEAT SHOCK PROTEIN.
 GN HSP30 OR YCR021C OR YCR21C.
 OS Saccharomyces cerevisiae (Baker's Yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
 NCBI_TaxID=4932;
 RN (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN=SKO2N;
 RX MEDLINE=93306747; PubMed=8319300;
 RA Regnacy M., Boucherie H.;
 RT "Isolation and sequence of HSP30, a yeast heat-shock gene coding for
 a hydrophobic membrane protein.";
 RL Curr. Genet. 23:435-442(1993).
 RN (2)
 RP SEQUENCE FROM N.A.
 RA Feldmann H., Manhaupt G., Vetter I.;
 RL Submitted (Mar-1992) to the EMBL/Genbank/DBJ databases.
 CC -1- FUNCTION: PROBABLY COOPERATES WITH OTHER HEAT SHOCK PROTEINS IN
 THE TRANSLLOCATION OF POLYPEPTIDES THROUGH MEMBRANES. IT MAY
 COUNTERACT THE ALTERING EFFECT OF HEAT SHOCK ON THE PLASMA
 MEMBRANE.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
 CC -1- DEVELOPMENTAL STAGE: EXPRESSED DURING THE ENTRY INTO STATIONARY
 PHASE RESULTING FROM GLUCOSE LIMITATION.
 CC -1- SIMILARITY: BELONGS TO THE ARCHAEL OPSIN FAMILY. HSP30
 SUBFAMILY.
 CC -----
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 or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M93123; AAA02903.1; -;
 DR EMBL; X59720; AAA02313.1; -;
 DR PIR; S30781; S30781.
 DR PIR; S19432; S19432.
 DR SGD; S0000615; HSP30.
 KW Heat shock; Transmembrane.
 FT TRANSMEM 35 55
 FT TRANSMEM 66 86 POTENTIAL.
 FT TRANSMEM 122 142 POTENTIAL.
 FT TRANSMEM 158 178 POTENTIAL.
 FT TRANSMEM 182 202 POTENTIAL.
 FT TRANSMEM 216 236 POTENTIAL.

FT TRANSMEM 249 269 POTENTIAL.
 FT DOMAIN 290 332 GLU-RICH (ACIDIC).
 SQ SEQUENCE 332 AA; 37044 MW; 260474A481D29AC5 CRC64;

Query Match 64.4%; Score 38; DB 1; Length 332;
 Best Local Similarity 87.5%; Pred. No. 16;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 HEAPEAPE 8
 DB 318 HEPEPEAE 325

RESULT 10
 DDN1_BOVIN STANDARD; PRT; 226 AA.
 AC P80219;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 15-DEC-1998 (Rel. 37, Last annotation update)
 DE DUODENASE I (EC 3.4.21.-) (DUODENUM SERINE PROTEASE).
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidea;
 OC Bovidae; Bovinae; Bos.
 NCBI_TaxID=9913;
 RN (1)
 RP SEQUENCE.
 RC TISSUE=Duoenum;
 RX MEDLINE=95172076; PubMed=7867649;
 RA Zamolodchikova T.S., Vorolytseva T.I., Nazimov I.V., Grishina G.A.;
 RT "Duodenase, a new serine protease of unusual specificity from bovine
 duodenal mucosa. Primary structure of the enzyme.";
 RL Eur. J. Biochem. 227:873-879(1995).
 RN (2)
 RP SEQUENCE OF 1-20 AND 172-183.
 RX MEDLINE=93048618; PubMed=1425193;
 RA Antonov V.K., Vorolytseva T.I., Zamolodchikova T.S.;
 RT "Duodenase -- a new serine proteinase with unusual specificity.";
 RL Dokl. Akad. Nauk SSSR 324:1318-1322(1992).
 RN (3)
 RP SEQUENCE OF 1-24.
 RX MEDLINE=95172075; PubMed=7867648;
 RA Zamolodchikova T.S., Vorolytseva T.I., Antonov V.K.;
 RT "Duodenase, a new serine protease of unusual specificity from bovine
 duodenal mucosa. Purification and properties.";
 RL Eur. J. Biochem. 227:866-872(1995).
 CC -1- CATALYTIC ACTIVITY: PREFERENTIAL CLEAVAGE: LYS-, ARG-, TYR-, PHE-,
 LEU-.
 CC -1- SUBUNIT: MONOMER.
 CC -1- MISCELLANEOUS: THE OPTIMUM PH AND TEMPERATURE OF DUODENASE I ARE
 8.0 AND 50 DEGREES CELSIUS, RESPECTIVELY.
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
 HSSP; P00763; IDPO.
 CC HSSP; P00763; IDPO.
 DR MEROPS; S01.142; -;
 DR InterPro; IPR001314; Chymotrypsin.
 DR InterPro; IPR001254; Trypsin.
 DR Pfam; PF00089; trypsin; 1.
 DR PRINTS; PR00722; CHYMOTRYPSIN.
 DR SMART; SM00020; TRY-SPC; 1.
 DR PROSITE; PS50240; TRYPSIN_DOM; 1.
 DR PROSITE; PS00134; TRYPSIN_HIS; 1.
 DR PROSITE; PS00135; TRYPSIN_SER; 1.
 KW Hydrolase; Serine protease; Glycoprotein.
 FT ACT_SITE 44 44
 FT ACT_SITE 87 87 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT ACT_SITE 181 181 CHARGE RELAY SYSTEM (BY SIMILARITY).
 FT DISULFD 29 45 BY SIMILARITY.
 FT DISULFD 121 187 BY SIMILARITY.
 FT DISULFD 152 166 BY SIMILARITY.
 FT CARBOHYD 50 50 N-LINKED (GLCNAC. . .).

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DR EMBL: J02907; AAA76859.1; -
DR EMBL: M57888; AAA03514.1; -
DR EMBL: M36118; AAA03248.1; -
DR EMBL: M72150; AAA74885.1; -
DR PIR: A32692; A32692.
DR HSSP: P04187; 2CP1.
DR MEROPS: S01.147; -.
DR MM: 116831; -.
DR InterPro: IPR001314; Chymotrypsin.
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; trypsin.1.
DR PRINTS: PR00722; CHYMOTRYPSIN.
DR SMART: SM00020; TRYP-Sec; 1.
DR PROSITE: PS0240; TRYP-SIN_DOM; 1.
DR PROSITE: PS00134; TRYP-SIN_HIS; 1.
DR PROSITE: PS00135; TRYP-SIN_SER; 1.
KW Hydrolase; Serine protease; Zymogen; Signal; T-cell; Cytolysis.
FT SIGNAL 1 18
FT PROPEP 19 20 ACTIVATION PEPTIDE.
FT CHAIN 21 246
FT ACT_SITE 64 246 GRANZYME H.
FT ACT_SITE 108 108 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 202 202 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT DISULFID 49 65 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT DISULFID 142 208 BY SIMILARITY.
FT DISULFID 172 187 BY SIMILARITY.
FT CARBOHYD 71 187 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 104 104 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 179 179 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 246 AA: 27315 MW: 6A6A873DA5F1E71 CRC64;

Query Match 62.7%; Score 37; DB 1; Length 246;
Best Local Similarity 54.5%; Pred. No. 18;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPETIM 11
| | | | |
Db 25 HEAKPHSRPYM 35

RESULT 13
GRAB_HUMAN
ID GRAB_HUMAN STANDARD; PRT: 247 AA.
AC P10144;
DT 01-MAR-1989 (Rel. 10; Created)
DT 01-MAR-1989 (Rel. 10; Last sequence update)
DT 20-AUG-2001 (Rel. 40; Last annotation update)
DE GRANZYME B PRECURSOR (EC 3.4.21.79) (T-CELL SERINE PROTEASE 1-3F)
DE (CYTOTOXIC T-LYMPHOCYTE PROTEINASE 2) (LYMPHOCYTE PROTEASE) (SECT)
DE (GRANZYME 2) (CATHEPSIN G-LIKE 1) (CTSG1) (CTLA-1) (FRAGMENTIN 2)
DE (HUMAN LYMPHOCYTE PROTEIN) (HLP) (C11).
GN GZMB OR CTLA1 OR GRB OR CGBP OR CCL1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87224164; PubMed=2953613;
RA Schmid J., Weissman C.;
RT "Induction of mRNA for a serine protease and a
RT beta-thromboglobulin-like protein in mitogen-stimulated human
RT leukocytes.";
RL J. Immunol. 139:250-256(1987).
RP SEQUENCE FROM N.A.
RX MEDLINE=88196184; PubMed=3258665;

RA Caputo A., Fahey D., Lloyd C., Vozab R., McCalins E., Rowe P.B.;
RT "Structure and differential mechanisms of regulation of
RT a serine esterase gene in activated human T lymphocytes.";
RL J. Biol. Chem. 263:6363-6369(1988).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=88320548; PubMed=3261871;
RA Trapani J.A., Klein J.L., White P.C., Dupont B.;
RT "Molecular cloning of an inducible serine esterase gene from human
RT cytotoxic lymphocytes.";
RL Proc. Natl. Acad. Sci. U.S.A. 85:6924-6928(1988).
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=89357968; PubMed=2788607;
RA Klein J.L., Shows T.B., Dupont B., Trapani J.A.;
RT "Genomic organization and chromosomal assignment for a serine
RT protease gene (CSPB) expressed by human cytotoxic lymphocytes.";
RL Genomics 5:110-117(1989).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=90308320; PubMed=2365998;
RA Caputo A., Sauer D.E., Rowe P.B.;
RT "Nucleotide sequence and genomic organization of a human T lymphocyte
RT serine protease gene.";
RL J. Immunol. 145:737-744(1990).
RN [6]
RP SEQUENCE FROM N.A.
RX MEDLINE=90236319; PubMed=2332171;
RA Haddad P., Clement M.V., Bernard O., Larsen C.J., Degos L.,
RA Sasportes M., Mathieu-Mahul D.;
RT "Structural organization of the CTLA-1 gene encoding human granzyme
RT B.";
RL Gene 87:265-271(1990).
RN [7]
RP SEQUENCE OF 21-40, AND CHARACTERIZATION.
RX MEDLINE=89009866; PubMed=3262682;
RA Hameed A., Lowrey D.M., Lichtenheld M., Podack E.R.;
RT "Characterization of three serine esterases isolated from human IL-2
RT activated killer cells.";
RL J. Immunol. 141:3142-3147(1988).
RN [8]
RP SEQUENCE OF 21-40, AND CHARACTERIZATION.
RX MEDLINE=89035468; PubMed=3263427;
RA Kraehenbuhl O., Rey C., Jenne D.E., Lanzavecchia A., Groscurth P.,
RA Carrel S., Tschopp J.;
RT "Characterization of granzymes A and B isolated from granules of
RT cloned human cytotoxic T lymphocytes.";
RL J. Immunol. 141:3471-3477(1988).
RN [9]
RP SEQUENCE OF 21-38.
RX MEDLINE=91093203; PubMed=1985927;
RA Poe M., Blake J.T., Boulton D.A., Gammon M., Sigal N.H., Wu J.K.,
RA Zweerink H.J.;
RT "Human cytotoxic lymphocyte granzyme B: its purification from
RT granules and the characterization of substrate and inhibitor
RT specificity.";
RL J. Biol. Chem. 266:98-103(1991).
RN [10]
RP FUNCTION: THIS ENZYME IS NECESSARY FOR TARGET CELL LYSIS IN CELL-
RP MEDIATED IMMUNE RESPONSES. IT CLEAVES AFTER ASP. SEEMS TO BE
RP LINKED TO AN ACTIVATION CASCADE OF CASPASES (ASPARTATE-SPECIFIC
RP CYSTEINE PROTEASES) RESPONSIBLE FOR APOPTOSIS EXECUTION. CLEAVES
RP CASPASE-3, -7, -9 AND 10 TO GIVE RISE TO ACTIVE ENZYMES MEDIATING
RP APOPTOSIS.
RN [11]
RP CATALYTIC ACTIVITY: PREFERENTIAL CLEAVAGE: ASP-I-XNA >> ASN-I-XNA
RN [12]
RP SUBCELLULAR LOCATION: CYTOPLASMIC GRANULES OF CYTOLYTIC T-
RN [13]
RP LYMPHOCYTES AND NATURAL KILLER CELLS.
RN [14]
RP INDUCTION: BY STAPHYLOCOCCAL ENTEROTOXIN A (SEA) IN PERIPHERAL
RN [15]
RP BLOOD LEUKOCYTES.
RN [16]
RP SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1: ALSO KNOWN AS THE
RN [17]
RP TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
RN [18]
RP PROTEASES.

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DR EMBL: M17016; AAA36627.1; -
 DR EMBL: J03189; AAA36603.1; -
 DR EMBL: J04071; AAA52118.1; -
 DR EMBL: J03072; AAB59528.1; -
 DR EMBL: M38193; AAA67124.1; -
 DR EMBL: M28879; AAA75490.1; -
 DR PIR: A28659; A28659; -
 DR PIR: A31405; A31405; -
 DR PIR: A32168; A32168; -
 DR PIR: B30525; B30525; -
 DR PIR: B30526; B30526; -
 DR HSSP: P04187; 2CP1; -
 DR MEROPS: S01.010; -
 DR MIM: I23910; -
 DR InterPro: IPR001254; Trypsin.
 DR Pfam: PF00089; trypsin; 1.
 DR SMART: SM00020; TRYP_SPE; 1.
 DR PROSITE: PS50240; TRYPSIN_DOM; 1.
 DR PROSITE: PS00134; TRYPSIN_HIS; 1.
 DR PROSITE: PS00135; TRYPSIN_SER; 1.
 DR Hydroxylase; Serine protease; Zymogen; Signal; T-cell; Cytolysis;
 KW Apoptosis.
 FT SIGNAL 1 18
 FT PROPEP 19 20
 FT CHAIN 21 247
 FT ACT_SITE 64 247
 FT ACT_SITE 108 108
 FT ACT_SITE 203 203
 FT DISULFID 49 65
 FT DISULFID 142 209
 FT DISULFID 173 188
 FT CARBOHYD 71 71
 FT CARBOHYD 104 104
 FT CONFLICT 55 55
 FT CONFLICT 72 72
 FT CONFLICT 94 94
 FT CONFLICT 212 212
 FT CONFLICT 247 AA; 27688 MM; 684FF605D6F2F4FB CRC64;
 SQ SEQUENCE

Query Match 62.7%; Score 37; DB 1; Length 247;
 Best Local Similarity 54.5%; Pred. No. 18;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
 |||:|:|
 Db 25 HEAKPHSRPYM 35

RESULT 14
 TONB_SERMA STANDARD; PRT; 247 AA.
 AC P26185;
 DT 01-MAY-1992 (Rel. 22, Created)
 DT 01-MAY-1992 (Rel. 22, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE TONB PROTEIN.
 GN TONB.
 OS Serratia marcescens.
 CC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 CC Serratia.
 CC NCBI_TaxID=615;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-W225;

RX MEDLINE=92140042; PubMed=1838128;
 RA Gaisser S., Braun V.;
 RT "The tonB gene of Serratia marcescens: sequence, activity and partial
 RT complementation of Escherichia coli tonB mutants.";
 RL Mol. Microbiol. 5:2777-2787 (1991).
 CC -1- FUNCTION: INTERACTS WITH OUTER MEMBRANE RECEPTOR PROTEINS THAT
 CC CARRY OUT HIGH-AFFINITY BINDING AND ENERGY DEPENDENT UPTAKE INTO
 CC THE PERIPLASMIC SPACE OF SPECIFIC SUBSTRATES SUCH AS COBALAMIN,
 CC AND VARIOUS IRON COMPOUNDS (SUCH AS IRON DICTYRATE, ENTEROCHELIN,
 CC AEROBACTIN, ETC.). IN THE ABSENCE OF TONB THESE RECEPTORS BIND
 CC THEIR SUBSTRATES BUT DO NOT CARRY OUT ACTIVE TRANSPORT. TONB ALSO
 CC INTERACTS WITH SOME COLICINS AND IS INVOLVED IN THE ENERGY-
 CC DEPENDENT, IRREVERSIBLE STEPS OF BACTERIOPHAGES PH1-80 AND T1
 CC INFECTION. IT COULD ACT TO TRANSDUCE ENERGY FROM THE CYTOPLASMIC
 CC MEMBRANE TO SPECIFIC ENERGY-REQUIRING PROCESSES IN THE OUTER
 CC MEMBRANE, RESULTING IN THE RELEASE INTO THE PERIPLASM OF LIGANDS
 CC BOUND BY THESE OUTER MEMBRANE PROTEINS (BT SIMILARITY).
 CC -1- SUBUNIT: THE ACCESSORY PROTEINS EXB8 AND EXB2 SEEM TO FORM A
 CC COMPLEX WITH TONB.
 CC -1- SUBCELLULAR LOCATION: PERIPLASMIC. ANCHORED TO THE CYTOPLASMIC
 CC MEMBRANE VIA ITS N-TERMINAL SIGNAL-LIKE SEQUENCE. SPANS THE
 CC PERIPLASM.
 CC -1- SIMILARITY: BELONGS TO THE TONB FAMILY.

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DR EMBL: X60996; GAA43308.1; -
 DR InterPro: IPR003538; TonB.
 DR Transport; Protein transport; Bacteriocin transport; Inner membrane;
 KW Periplasmic; Transmembrane; Signal-anchor; Repeat; Phage recognition.
 FT DOMAIN 1 10
 FT TRANSMEM 12 35
 FT DOMAIN 36 247
 FT DOMAIN 76 85
 FT DOMAIN 101 110
 FT DOMAIN 247 AA; 27389 MM; 46EBE6869EDB864B CRC64;
 SQ SEQUENCE

Query Match 62.7%; Score 37; DB 1; Length 247;
 Best Local Similarity 70.0%; Pred. No. 18;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 EAPEPEPEPIM 11
 |||||
 Db 78 EPEPEPEPIV 87

RESULT 15
 NKPL_RAT STANDARD; PRT; 248 AA.
 AC P18291;
 DT 01-NOV-1990 (Rel. 16, Created)
 DT 01-NOV-1990 (Rel. 16, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE NATURAL KILLER CELL PROTEASE 1 PRECURSOR (EC 3.4.21.-) (RNKP-1)
 DE (FRAGMENTIN).
 OS Rattus norvegicus (Rat).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 CC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RP TISSUE=T-cell;
 RX MEDLINE=90171602; PubMed=2307850;
 RA Zunino S.J., Bleackley R.C., Martinez J., Hudig D.;
 RT "RNKP-1, a novel natural killer killer-associated serine protease gene
 RT cloned from RNK-16 cytotoxic lymphocytes.";

RL J. Immunol. 144:2001-2009(1990).
RN [2]
RP SEQUENCE OF 21-53.
RX MEDLINE-92091788; PubMed=1727874;
RA Sayers T.J., Wiltrout T.A., Sowder R., Munger W.L., Smyth M.J.,
RA Henderson L.E.;
RT "Purification of a factor from the granules of a rat natural killer
RT cell line (RNK) that reduces tumor cell growth and changes tumor
RT morphology. Molecular identity with a granule serine protease
RT (RNKP-1).";
RL J. Immunol. 148:292-300(1992).
RN [3]
RP PARTIAL SEQUENCE.
RX MEDLINE-92121838; PubMed=1732416;
RA Shi L., Kraut R.P., Aebersold R., Greenberg A.H.;
RT "A natural killer cell granule protein that induces DNA fragmentation
RT and apoptosis.";
RL J. Exp. Med. 175:553-566(1992).
CC -1- FUNCTION: THIS ENZYME IS PROBABLY NECESSARY FOR TARGET CELL
CC LYSIS IN CELL-MEDIATED IMMUNE RESPONSES.
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S1; ALSO KNOWN AS THE
CC TRYPSIN FAMILY. STRONGEST TO OTHER GRANZYMES AND TO MAST CELL
CC PROTEASES.
CC -----
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CC or send an email to license@isb-stb.ch).
CC -----
DR EMBL: M34097; AAA42055.1; -.
DR PIR: A43520; A43520.
DR HSSP: P04187; 2CPI.
DR MEROPS: S01.136; -.
DR InterPro: IPR001254; Trypsin.
DR Pfam: PF00089; Trypsin.1.
DR SMART: SM00020; TRYP_SPC; 1.
DR PROSITE: PS50240; TRYP_SIN_DOM; 1.
DR PROSITE: PS00134; TRYP_SIN_HIS; 1.
DR PROSITE: PS00135; TRYP_SIN_SER; 1.
KW Hydrolase; Serine protease; Zymogen; Signal; T-cell; Cytolysis.
KW SIGNAL
FT 1 18
FT PROPEP 19 20 ACTIVATION PEPTIDE.
FT CHAIN 21 248 NATURAL KILLER CELL. PROTEASE 1.
FT ACT_SITE 65 65 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 109 109 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT ACT_SITE 204 204 CHARGE RELAY SYSTEM (BY SIMILARITY).
FT DISULFID 50 66 BY SIMILARITY.
FT DISULFID 143 210 BY SIMILARITY.
FT DISULFID 174 189 BY SIMILARITY.
FT DISULFID 98 98 H -> F (IN REF. 3).
FT CONFLICT 138 138 K -> D (IN REF. 3).
SQ SEQUENCE 248 AA; 27326 MW; 6F52089DDACC88C CRC64;

Query Match 62.7%; Score 37; DB 1; Length 248;
Best Local Similarity 54.5%; Pred. No. 18;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
| | | | : | |
DB 25 HEAKPHSRPYM 35

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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:42 ; Search time 27.18 seconds
(without alignments)
30.829 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAPEAPEIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues
Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR-68:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	42	71.2	144	2 S35331	apidaecin 22 precu
2	42	71.2	283	2 S35332	apidaecin 73 precu
3	40	67.8	429	2 S34224	cyellin B - long-ta
4	39	66.1	251	2 T39332	hypothetical prote
5	39	66.1	486	2 S07633	hematopoietic line
6	39	66.1	848	2 T23694	hypothetical prote
7	39	66.1	1593	2 T22028	hypothetical prote
8	38	64.4	150	2 C72724	hypothetical prote
9	38	64.4	168	2 S35330	apidaecin 14 precu
10	38	64.4	248	2 S33755	granzyme-like prot
11	38	64.4	256	1 PRH03	apidaecin 3 (EC 3
12	38	64.4	288	2 S1059	proteinsase 3 (EC 3
13	38	64.4	332	2 S31848	hypothetical prote
14	37	62.7	21	2 S69371	heat shock protein
15	37	62.7	125	2 T36367	duodenase - bovine
16	37	62.7	175	2 T05669	hypothetical prote
17	37	62.7	226	2 S69370	hypothetical prote
18	37	62.7	246	2 A32692	duodenase - bovine
19	37	62.7	247	2 S18592	cytotoxic T-lympho
20	37	62.7	248	2 S43259	tonB protein - Ser
21	37	62.7	248	2 A43520	granzyme-like prot
22	37	62.7	251	2 T10262	natural killer cel
23	37	62.7	265	2 B81138	mast cell serine p
24	37	62.7	265	2 C81883	phosphatidylserine
25	37	62.7	269	2 E69381	probable membrane
26	37	62.7	281	1 A61021	hypothetical prote
27	37	62.7	356	2 T00881	granzyme B (EC 3.4
28	37	62.7	376	2 S55892	probable PCP2-like
29	37	62.7	464	2 S00757	endo-1,4-beta-xyla
					deoxyribodipyrimid

30	37	62.7	612	2 T42243	probable polypepti
31	37	62.7	1280	2 T00365	hypothetical prote
32	37	62.7	2109	2 T33247	hypothetical prote
33	37	62.7	2584	2 T24158	hypothetical prote
34	37	62.7	2606	2 T24157	hypothetical prote
35	37	62.7	3119	2 T18414	protein 9377 - mal
36	37	62.7	5138	2 B96695	hypothetical prote
37	36	61.0	182	2 T16423	hypothetical prote
38	36	61.0	206	2 B48441	antigen (C-termina
39	36	61.0	239	1 BVEC	tonB protein - Esc
40	36	61.0	239	2 G85705	hypothetical prote
41	36	61.0	240	2 S13257	tonB protein - Sal
42	36	61.0	243	2 T45505	membrane protein
43	36	61.0	255	2 S30280	tonB protein - Yer
44	36	61.0	265	2 T00765	hypothetical prote
45	36	61.0	266	2 T44781	tonB protein (impo

ALIGNMENTS

RESULT 1
S35331
apidaecin 22 precursor - honeybee
C:Species: Apis mellifera (honeybee)
C>Date: 03-Feb-1994 #sequence-revision 03-Feb-1994 #text-change 21-Jul-2000
C:Accession: S35331
R:Castels-Josson, K.; Capaci, T.; Castels, P.; Tempst, P.
EMBO J. 12, 1569-1578, 1993
A>Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplifi
A:Reference number: S35330; MUID:93223697
A:Accession: S35331
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-144 <CAS>
A:Cross-references: EMBL:X72576; NID:9297064; PIDN:CAAS1168.1; PID:9297065
C:Superfamily: procyclic acidic repetitive protein

Query Match 71.2%; Score 42; DB 2; Length 144;
Best Local Similarity 100.0%; Pred. No. 2.2;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 EAPEAP 9
DB 35 EAPEAP 42

RESULT 2
S35332
apidaecin 73 precursor - honeybee (fragment)
N:Contains: apidaecin 1a
C:Species: Apis mellifera (honeybee)
C>Date: 03-Feb-1994 #sequence-revision 03-Feb-1994 #text-change 03-Nov-2000
C:Accession: S35332; S05383
R:Castels-Josson, K.; Capaci, T.; Castels, P.; Tempst, P.
EMBO J. 12, 1569-1578, 1993
A>Title: Apidaecin multipeptide precursor structure: a putative mechanism for amplifi
A:Reference number: S35330; MUID:93223697
A:Accession: S35332
A:Molecule type: mRNA
A:Residues: 1-283 <CAS>
A:Cross-references: EMBL:X72577; NID:9297066; PIDN:CAAS1169.1; PID:94539289
A:Accession: S05383
A:Molecule type: protein
A:Residues: 258-283 <CAS>
C:Superfamily: proline-rich protein
F:266-283/Product: apidaecin 1a #status experimental <MAT>

Query Match 71.2%; Score 42; DB 2; Length 283;
Best Local Similarity 100.0%; Pred. No. 4.5;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
| | | | |
Db 62 EAEPEAP 69

RESULT 3
S34224
cyclin B - long-tailed hamster

C:Species: Cricetus longicaudatus (long-tailed hamster)

C>Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 16-Jul-1999

C:Accession: S34224

R:Markiewicz, D.A.; Flick, M.B.; Mushel, R.J.; McKenna, W.G.

submitted to the EMBL Data Library, March 1992

A:Description: New features of mammalian cyclins seen in rat and chinese hamster cyclin

A:Reference number: S34224

A:Accession: S34224

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-429 <MAR>

A:Cross-references: EMBL:X64588; NID:g313764; PIDN:CAA45876.1; PID:g313765

C:Superfamily: cyclin

C:Keywords: cell cycle control

Query Match 67.8%; Score 40; DB 2; Length 429;
Best Local Similarity 70.0%; Pred. No. 16;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 2 EAEPEAP 11
| | | | |
Db 98 EPEPEPEPV 107

RESULT 4

T39332
hypothetical protein SPBC11G11.05 - fission yeast (Schizosaccharomyces pombe)

C:Species: Schizosaccharomyces pombe

C>Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999

C:Accession: T39332

R:Saunders, D.; Harris, D.; McDougall, R.C.; Rajandream, M.A.; Barrell, B.G.

submitted to the EMBL Data Library, October 1999

A:Reference number: T39332

A:Accession: T39332

A>Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-251 <SAU>

A:Cross-references: EMBL:AL132747; PIDN:CA859807.1; GSPDB:GN00067; SPDB:SPBC11G11.05

A:Experimental source: strain 972h; cosmid c11G11

C:Genetics:

A:Gene: SPDB:SPBC11G11.05

A:Map position: 2

Query Match 66.1%; Score 39; DB 2; Length 251;
Best Local Similarity 87.5%; Pred. No. 13;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 EAEPEAP 9
| | | | |
Db 185 ESEPEAP 192

RESULT 5

S07633
hematopoietic lineage cell-specific protein HSL - human

C:Species: Homo sapiens (man)

C>Date: 29-Jan-1993 #sequence_revision 29-Jan-1993 #text_change 05-Nov-1999

C:Accession: S07633; A47478; B47478; C47478; D47478; E47478

R:Klimmura, D.; Kaneko, H.; Miyagoe, Y.; Aiyasu, T.; Watanabe, T.

Nucleic Acids Res. 17, 9367-9379, 1989

A:Title: Isolation and characterization of a novel human gene expressed specifically in

A:Reference number: S07633; M0ID:90067934

A:Accession: S07633

A:Molecule type: mRNA

A:Residues: 1-486 <KIT>

A:Cross-references: EMBL:X16663; NID:g32054; PIDN:CAA34651.1; PID:g32055

R:Yamanashi, Y.; Okada, M.; Sema, T.; Yamori, T.; Umemori, H.; Tsunashima, S.; Toyosh

Proc. Natl. Acad. Sci. U.S.A. 90, 3631-3635, 1993

A:Title: Identification of HSL protein as a major substrate of protein-tyrosine kinases

A:Reference number: A47478; M0ID:93234551

A:Accession: A47478

A>Status: preliminary

A:Molecule type: protein

A:Residues: 4-19, 'XXX', 23-26, 'X', 79-93, 'X', 95, 'X', 134-146, 208-223, 274-284, 'X', 286, 'X'

A:Experimental source: Dauid, B-lymphoblastoid cells

A>Note: sequence modified after extraction from NCBI backbone

C:Superfamily: SH3 homology

F:435-482/Domain: SH3 homology <SH3>

Query Match 66.1%; Score 39; DB 2; Length 486;
Best Local Similarity 77.8%; Pred. No. 27;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 HEAEPEAP 9
| | | | |
Db 360 YEAEPEPEP 368

RESULT 6

T23694
hypothetical protein M03C11.2 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

C:Accession: T23694

R:McMurray, A.

submitted to the EMBL Data Library, April 1995

A:Reference number: T23694

A:Accession: T23694

A>Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-848 <MIL>

A:Cross-references: EMBL:Z49128; PIDN:CA88959.1; GSPDB:GN00021; CESP:M03C11.2

A:Experimental source: clone M03C11

C:Genetics:

A:Gene: CESP:M03C11.2

A:Map position: 3

A:Introns: 113/2; 147/3; 185/3; 379/1; 482/3; 553/2; 688/3; 762/3

Query Match 66.1%; Score 39; DB 2; Length 848;
Best Local Similarity 77.8%; Pred. No. 48;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAEPEAP 10
| | | | |
Db 506 EPEPEAP 514

RESULT 7

T22028
hypothetical protein F40F11.2 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999

C:Accession: T22028

R:Dobson, R.

submitted to the EMBL Data Library, May 1996

A:Reference number: Z19504

A:Accession: T22028

A>Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-1593 <MIL>

A:Cross-references: EMBL:Z73426; NID:el343215; PIDN:CAA97793.1; GSPDB:GN00022; CESP:F

A:Experimental source: clone F40F11

C:Genetics:

A:Reference number: JH0331; MUID:91079774
 A:Accession: JH0331
 A:Molecule type: mRNA
 A:Residues: 'R',3-118, 'V',120-134, 'MT',137-254, 'P' <CAM>
 A:Cross-references: GB:X55668; NID:935687; PIDN:CAA39203.1; PID:9135280
 A:Note: part of this sequence, including the amino end of the mature protein, was confir
 R:Bories, D.; Raynal, M.C.; Solomon, D.H.; Darzynkiewicz, Z.; Cayre, Y.E.
 Cell 59, 959-968, 1989
 A:Title: Down-regulation of a serine protease, myeloblastin, causes growth arrest and d
 A:Reference number: A33751; MUID:90090622
 A:Accession: A33751
 A:Molecule type: mRNA
 A:Residues: 42-256 <BO>
 A:Cross-references: GB:M29142; NID:9188983; PIDN:AA36342.1; PID:9188984
 A:Note: the authors translated the codon GGG for residue 49 as G10, GGC for residue 52 a
 R:Jenne, D.E.; Tschopp, J.; Luedemann, J.; Utecht, B.; Gross, W.L.
 Nature 346, 520, 1990
 A:Title: Wegener's antineutrophil proteinase.
 A:Reference number: S11091; MUID:90332035
 A:Accession: S11091
 A:Molecule type: mRNA
 A:Residues: 20-56 <JEN>
 R:Musette, P.; Labbaye, C.; Dorner, M.H.; Cayre, Y.E.; Casanova, J.L.; Kourilsky, P.
 Blood 77, 1398-1399, 1991
 A:Title: Wegener's antineutrophil proteinase and leukemia.
 A:Reference number: A61176; MUID:91159650
 A:Accession: A61176
 A:Molecule type: mRNA
 A:Residues: 1-42 <MUS>
 A:Cross-references: EMBL:X56606; NID:935189; PIDN:CAA39943.1; PID:935190
 APMIS 19(Suppl.), 26-27, 1990
 R:Goldschmeding, R.; Dolman, K.M.; Van Den Ende, M.E.; Van Der Meer-Gerritsen, C.H.; So
 A:Title: The relation of 29 kD C-ANCA antigen to proteinase 3.
 A:Reference number: A60148; MUID:91136884
 A:Accession: A60148
 A:Molecule type: protein
 A:Residues: 28-48 <GOL>
 R:Rao, N.V.; Wehner, N.G.; Marshall, B.C.; Gray, W.R.; Gray, B.H.; Hoidal, J.R.
 J. Biol. Chem. 266, 9540-9548, 1991
 A:Title: Characterization of proteinase-3 (PR-3), a neutrophil serine proteinase. Struct
 A:Reference number: A43982; MUID:91236723
 A:Accession: A43982
 A:Molecule type: protein
 A:Residues: 28-61, 'X',63, 'D',65-67,228-244 <RAO>
 R:Wilde, C.G.; Snable, J.L.; Griffith, J.E.; Scott, R.W.
 J. Biol. Chem. 265, 2038-2041, 1990
 A:Title: Characterization of two azurophil granule proteases with active-site homology t
 A:Reference number: A43981; MUID:90130450
 A:Accession: A43981
 A:Molecule type: protein
 A:Residues: 28-45, 'E',47,196-208, 'X',210-215, 'X',217-219 <WIL>
 R:Gabay, J.E.; Scott, R.W.; Campanelli, D.; Griffith, J.; Wilde, C.; Marra, M.N.; Seeger
 Proc. Natl. Acad. Sci. U.S.A. 86, 5610-5614, 1989
 A:Title: Antibiologic proteins of human polymorphonuclear leukocytes.
 A:Reference number: A33913; MUID:89315847
 A:Accession: C33913
 A:Molecule type: protein
 A:Residues: 28-45, 'E',47 <GAB>
 R:Niles, J.L.; McCluskey, R.T.; Almad, M.F.; Arnaout, M.A.
 Blood 74, 1888-1893, 1989
 A:Title: Wegener's granulomatosis antineutrophil proteinase is a novel neutrophil serine proteinase.
 A:Reference number: A60481; MUID:90028708
 A:Accession: A60481
 A:Molecule type: protein
 A:Residues: 28-38, 'X',40-47 <NIL>
 R:Ohlsson, K.; Linder, C.; Rosenzweig, M.
 Biol. Chem. Hoppe-Seyler 371, 549-555, 1990
 A:Title: Monoclonal antibodies specific for neutrophil proteinase 4. Production and use
 A:Reference number: S10605; MUID:91025622
 A:Accession: S10605
 A:Molecule type: protein
 A:Residues: 28-52 <OHL>
 R:Luedemann, J.; Utecht, B.; Gross, W.L.

J. Exp. Med. 171, 357-362, 1990
 A:Title: Anti-neutrophil cytoplasm antibodies in Wegener's granulomatosis recognize a
 A:Reference number: PL0230; MUID:90111630
 A:Accession: PL0230
 A:Molecule type: protein
 A:Residues: 28-37, 'I',39-40, 'I',41-43 <LUE>
 C:Comment: This polymorphonuclear leukocyte serine protease from azurophilic granules
 C:Genetics:
 A:Gene: GDB:PRN3
 A:Cross-references: GDB:126876; OMIM:177020
 A:Map position: 19p13.3-19p13.3
 A:Introns: 21/L; 76/2; 123/3; 200/3
 C:Superfamily: trypsin; trypsin homology
 C:Keywords: glycoprotein; hydrolase; polymorphonuclear leukocyte; serine proteinase
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 F:26-27/Domain: propeptide #status predicted <PRO>
 F:28-256/Product: proteinase 3 #status experimental <MNT>
 F:28-243/Domain: trypsin homology <TRY>
 F:56-72,152-209,182-188,199-224/Disulfide bonds: #status predicted
 F:71,118,203/Active site: His, Asp, Ser #status predicted
 F:129,174/Binding site: carboxylate (Asn) (covalent) #status predicted

Query Match	Score 38;	DB 1;	Length 256;
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			Indels 0;
			Gaps 0;

QY 1 HEAPEPEPTM 11
 DB 32 HEAPEPEPTM 42

RESULT 12
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 hypothetical protein B12F1.110 [imported] - Neurospora crassa
 C:Species: Neurospora crassa
 C:Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
 C:Accession: T51059
 R:Schlatter, U.; Aign, V.; Hohnsels, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatu
 submitted to the Protein Sequence Database, July 2000
 A:Reference number: Z25286
 A:Accession: T51059
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-288 <SCH>
 A:Cross-references: EMBL:AL390091; GSPDB:GN00116; NCSP:B12F1.110
 A:Experimental source: BAC clone B12F1, strain OR74A
 C:Genetics:
 A:Gene: NCSP:B12F1.110
 A:Map position: 6
 A:Introns: 154/1

Query Match	Score 38;	DB 2;	Length 288;
Best Local Similarity	54.5%;	Pred. No. 23;	
Matches 6;	Conservative	2;	Mismatches 3;
			Indels 0;
			Gaps 0;

QY 1 HEAPEPEPTM 11
 DB 58 HEAPEPEPTM 68

RESULT 13
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 heat shock protein HSP30 - yeast (Saccharomyces cerevisiae)
 N:Alternate names: protein YCR021c
 C:Species: Saccharomyces cerevisiae
 C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 29-Oct-1999
 C:Accession: S31848; S19432; S30781
 R:Regnacy, M.; Boucherie, H.
 Curr. Genet. 23, 435-442, 1993
 A:Title: Isolation and sequence of HSP30, a yeast heat-shock gene coding for a hydrop
 A:Reference number: S31848; MUID:93306747

A:Accession: S31848
 A:Molecule type: DNA
 A:Residues: 1-263 <REG1>
 A:Cross-references: EMBL:M93123
 R:Feldmann, H.; Mannhaupt, G.; Vetter, I.
 submitted to the Protein Sequence Database, March 1992
 A:Reference number: S19429
 A:Accession: S19432
 A:Molecule type: DNA
 A:Residues: 1-190, 'A', 192-332 <FEU>
 A:Cross-references: EMBL:X59720; NID:G1907116; PIDN:CAA42313.1; PID:e264485; PID:G190716
 R:Regnacy, M.; Boucherie, H.
 submitted to the EMBL Data Library, January 1993
 A:Reference number: S30781
 A:Accession: S30781
 A:Molecule type: DNA
 A:Residues: 1-152, 'GY', 164-165, 167, 'A', 169, 'NSNRGL', 170-240, 'VFNQT', 270, 278-280, 'FNVFVH'
 A:Cross-references: EMBL:M93123
 A:Note: the difference at the carboxyl end is due to a frameshift error
 C:Genetics:
 A:Gene: SGD:HSP30
 A:Cross-references: SGD:S0000615; MIPS:YCR021C
 A:Map position: 3R
 C:Keywords: membrane protein

Query Match 64.4%; Score 38; DB 2; Length 332;
 Best Local Similarity 87.5%; Pred. No. 27;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 HEAPEAE 8
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 Db 318 HEPEAE 325

RESULT 14
 S69371
 duodenase - bovine (fragment)
 C:Species: Bos primigenius taurus (cattle)
 C:Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 21-Nov-1998
 C:Accession: S69371
 R:Zamolodchikova, T.S.; Vorolyntseva, T.I.; Antonov, V.K.
 Eur. J. Biochem. 227, 866-872, 1995
 A:Title: Duodenase, a new serine protease of unusual specificity from bovine duodenal mu
 A:Reference number: S69371; MUID:55172075
 A:Accession: S69371
 A:Molecule type: protein
 A:Residues: 1-21 <ZAK>
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Query Match 62.7%; Score 37; DB 2; Length 21;
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 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAEPIIM 11
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 Db 5 HEAKPHSRPYM 15

RESULT 15
 T36257
 hypothetical protein SCE68.07c - Streptomyces coelicolor
 C:Species: Streptomyces coelicolor
 C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999
 C:Accession: T36257
 R:Murphy, L.; Harris, D.; James, K.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
 submitted to the EMBL Data Library, June 1999
 A:Reference number: Z21576
 A:Accession: T36257
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-125 <MUR>

A:Cross-references: EMBL:AL079345; PIDN:CA945343.1; GSPDB:GND00070; SCOEDB:SCE68.07c
 A:Experimental source: strain A3(2)
 C:Genetics:
 A:Gene: SCOEDB:SCE68.07c

Query Match 62.7%; Score 37; DB 2; Length 125;
 Best Local Similarity 87.5%; Pred. No. 15;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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 Db 5 EAPEEAP 12

Search completed: January 4, 2002, 08:41:31
 Job time: 169 sec

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; Sequence 1, Application US/08479537A
; Patent No. 5861381
; GENERAL INFORMATION:
; APPLICANT: CHAMON, Pierre
; APPLICANT: KIENY, Marie-Paule
; APPLICANT: LATHE, Richard
; APPLICANT: HAREUVENT, Mara
; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
; TITILE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/479,537A
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 90/13101
; FILING DATE: 23-OCT-1990
; APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/FR91/00835
; FILING DATE: 23-OCT-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/039,320
; FILING DATE: 04-APR-1993
; APPLICATION DATA:
; APPLICATION NUMBER: US 08/403,576
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Teskin, Robin L.
; REGISTRATION NUMBER: 35,030
; REFERENCE/DOCKET NUMBER: 017753-025
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 6192 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: sig_peptide
; LOCATION: 58..120
; FEATURE:

NAME/KEY: repeat_region
LOCATION: 439..5239
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OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat i
OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fi
OTHER INFORMATION: The number of such repeats varies from 1 to 80."
NAME/KEY: mat_peptide
LOCATION: 121..6166
FEATURE:
NAME/KEY: repeat_region
LOCATION: 457
OTHER INFORMATION: /note= "Nucleotide 457 is X1 = NNN
OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC,
OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
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NAME/KEY: repeat_region
LOCATION: 487
OTHER INFORMATION: /note= "Nucleotide 487 is Y = NNN
OTHER INFORMATION: which is the codon for Thr or Asn wherein Thr = ACT, ACC,
OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
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OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
US-08-479-537A-1

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2 GUAlagluProgluAlagluPro 9
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seq_documentation_block:
; Sequence 1, Application US/09083116
; Patent No. 6203795
; GENERAL INFORMATION:
; APPLICANT: CHAMON, Pierre
; APPLICANT: KIENY, Marie-Paule
; APPLICANT: LATHE, Richard
; APPLICANT: HAREUVENT, Mara
; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
; TITILE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/083,116
; FILING DATE:
; CLASSIFICATION:
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OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
FEATURE:
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LOCATION: 487
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OTHER INFORMATION: which is the codon for Thr or Asn wherein Thr = ACT, ACC, ACN
OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
FEATURE:
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LOCATION: 496
OTHER INFORMATION: /note= "Nucleotide 496 is X2 = NNN
OTHER INFORMATION: which is the codon for Pro or Ala wherein Pro = CCT, CCC, CCA
OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
US-08-479-537A-4

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; Sequence 4, Application US/09083116
; Patent No. 6203795
; GENERAL INFORMATION:
; APPLICANT: CHAMBER, Pierre
; APPLICANT: KIENY, Marie-Paule
; APPLICANT: LAFITE, Richard
; APPLICANT: HAREVEU, Maria
; TITLE OF INVENTION: PHARMACEUTICAL COMPOSITION FOR THE
; TITRE OF INVENTION: TREATMENT OR PREVENTION OF A MALIGNANT TUMOR
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BURNS, DOANE, SWECKER & MATHIS, L.L.P.
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22313-1404
; COMPUTER READABLE FORM:
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; CURRENT APPLICATION DATA:
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; FILING DATE:
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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/479,537
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/FR91/00835
; FILING DATE: 23-OCT-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/039,320
; FILING DATE: 04-APR-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/403,576
; FILING DATE: 14-MAR-1995
; ATTORNEY/AGENT INFORMATION:

NAME: Teskin, Robin L.
REGISTRATION NUMBER: 35,030
REFERENCE/DOCKET NUMBER: 017753-025
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 836-6620
TELEFAX: (703) 836-2021
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 6449 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
FEATURE:
NAME/KEY: sig_peptide
LOCATION: 58..120
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NAME/KEY: repeat_region
LOCATION: 439..5239
OTHER INFORMATION: /note= "The nucleotides spanning
OTHER INFORMATION: 439-5239 constitute a repeated region wherein the repeat 1
OTHER INFORMATION: nucleotides and encodes 20 amino acids, 17 of which are fl
OTHER INFORMATION: The number of such repeats varies from 1 to 80."
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OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
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LOCATION: 487
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OTHER INFORMATION: or ACG; and Asn = AAT or AAC."
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LOCATION: 496
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OTHER INFORMATION: or CCG; and Ala = GCT, GCC, GCA, or GCG."
US-09-083-116-4

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seq_documentation_block:
; Sequence 12, Application US/08394600B
; Patent No. 5843693
; GENERAL INFORMATION:
; APPLICANT: Halenbeck, Robert F.
; APPLICANT: Jewell, David A.
; APPLICANT: Koths, Kirsten E.
; APPLICANT: Krieglert, Michael
; APPLICANT: Perez, Carl
; TITLE OF INVENTION: Compositions for the inhibition of

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: McAndrews, Held & Malloy, Ltd.
STREET: 500 West Madison Street, 34th Floor
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60661
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/394,600B
FILING DATE: 02/27/95
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Donald J. Pochoplen
REGISTRATION NUMBER: 32,167
REFERENCE/DOCKET NUMBER: 820.005/11850US05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/707-8889
TELEFAX: 312/707-9155
TELEX:
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 83 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
US-08-394-600B-12

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Percent Similarity: 90.909 Percent Identity: 54.545

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Sequence 12, Application PC/TUS9502513

GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

APPLICANT: Krieglner, Michael

APPLICANT: Perez, Carl

TITLE OF INVENTION: Compositions for the Inhibition of

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof

NUMBER OF SEQUENCES: 22

CORRESPONDENCE ADDRESS:

ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &

ADDRESS: Borun

STREET: 6300 Sears Tower, 233 South Wacker Drive

CITY: Chicago

STATE: Illinois

COUNTRY: United States of America

ZIP: 60606-6402

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 83 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA
PCT-US95-02513-12

alignment_scores:
Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-12/rev ..

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Sequence 3, Application US/08394600B

Patent No. 5843693

GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.

APPLICANT: Jewell, David A.

APPLICANT: Koths, Kirston E.

APPLICANT: Krieglner, Michael

APPLICANT: Perez, Carl

TITLE OF INVENTION: Compositions for the Inhibition of

TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof

NUMBER OF SEQUENCES: 28

CORRESPONDENCE ADDRESS:

ADDRESSEE: McAndrews, Held & Malloy, Ltd.

STREET: 500 West Madison Street, 34th Floor

CITY: Chicago

STATE: Illinois

COUNTRY: United States of America

ZIP: 60661

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/394,600B

FILING DATE: 02/27/95

CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:

NAME: Donald J. Pochoplen

REGISTRATION NUMBER: 32,167

REFERENCE/DOCKET NUMBER: 820.005/11850US05

TELECOMMUNICATION INFORMATION:
 TELEPHONE: 312/707-8889
 TELEFAX: 312/707-9155
 TELEEX:
 INFORMATION FOR SEQ ID NO: 3:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 771 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: cDNA
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1..771
 US-08-394-600B-3

alignment_scores:

Quality: 38.00 Length: 11
 Ratio: 3.800 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x US-08-394-600B-3 ..

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seq_name: /cgn2_6/ptodata/2/ina/5B_COMB.seq:US-08-394-600B-22

seq_documentation_block:

Sequence 22, Application US/08394600B
 Patent No. 5843693
 GENERAL INFORMATION:
 APPLICANT: Halenbeck, Robert F.
 APPLICANT: Jewell, David A.
 APPLICANT: Koths, Kirston E.
 APPLICANT: Krieglner, Michael
 APPLICANT: Perez, Carl
 TITLE OF INVENTION: Compositions for the inhibition of
 TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
 NUMBER OF SEQUENCES: 28
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: McAndrews, Heid & Malloy, Ltd.
 STREET: 500 West Madison Street, 34th Floor
 CITY: Chicago
 STATE: Illinois
 COUNTRY: United States of America
 ZIP: 60661
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/394,600B
 FILING DATE: 02/27/95
 CLASSIFICATION: 435
 ATTORNEY/AGENT INFORMATION:
 NAME: Donald J. Pochopien
 REGISTRATION NUMBER: 32,167
 REFERENCE/DOCKET NUMBER: 820,005/11850US05
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 312/707-8889
 TELEFAX: 312/707-9155
 TELEEX:
 INFORMATION FOR SEQ ID NO: 22:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 771 base pairs
 TYPE: nucleic acid

STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: cDNA
 FEATURE:
 NAME/KEY: CDS
 LOCATION: 1..771
 US-08-394-600B-22

alignment_scores:

Quality: 38.00 Length: 11
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alignment_block:

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seq_documentation_block:

Sequence 3, Application US/08230428B
 Patent No. 5998378
 GENERAL INFORMATION:
 APPLICANT: Krieglner, Michael
 APPLICANT: Halenbeck, Robert F.
 APPLICANT: Perez, Carl
 APPLICANT: Jewell, David A.
 APPLICANT: Koths, Kirston E.
 TITLE OF INVENTION: Compositions for the inhibition of TNF
 TITLE OF INVENTION: Hormone Formation And Uses Thereof (As Amended)
 NUMBER OF SEQUENCES: 14
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: CHIRON CORPORATION Intellectual Property - R440
 STREET: 4560 Horton Street, P.O. Box 8097
 CITY: Emeryville
 STATE: California
 COUNTRY: United States of America
 ZIP: 94662-8097
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/230,428B
 FILING DATE: 19-APR-1994
 CLASSIFICATION: 514
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 07/905,546
 FILING DATE: 25-JUN-1992
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 07/395,253
 FILING DATE: 16-AUG-1989
 ATTORNEY/AGENT INFORMATION:
 NAME: Savereide, Paul B.
 REGISTRATION NUMBER: 36,914
 REFERENCE/DOCKET NUMBER: 0820,004
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (510) 601-2718
 TELEFAX: (510) 655-3542
 INFORMATION FOR SEQ ID NO: 3:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 771 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA

FEATURE:
NAME/KEY: CDS
LOCATION: 1..768
US-08-230-428B-3

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Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

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Align seg 1/1 to: US-08-230-428B-3 from: 1 to: 771

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Sequence 3, Application PC/TUS9502513
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kolts, Kirston E.
APPLICANT: Krieglner, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the Inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
PCT-US95-02513-3

alignment_scores:

Quality: 38.00 Length: 11

Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-3

Align seg 1/1 to: PCT-US95-02513-3 from: 1 to: 771

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seq_name: /cgn2_6/prodata/2/ina/PCTUS_COMB.seq:PCT-US95-02513-22

seq_documentation_block:

Sequence 22, Application PC/TUS9502513
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kolts, Kirston E.
APPLICANT: Krieglner, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the Inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6300
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 22:
SEQUENCE CHARACTERISTICS:
LENGTH: 771 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..771
PCT-US95-02513-22

alignment_scores:

Quality: 38.00 Length: 11
Ratio: 3.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 54.545

alignment_block:

US-09-444-281-27 x PCT-US95-02513-22

Align seg 1/1 to: PCT-US95-02513-22 from: 1 to: 771

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seq_documentation_block:
; Sequence 645, Application US/08974549A
; Patent No. 6166178
; GENERAL INFORMATION:
; APPLICANT: Cech, Thomas R.
; APPLICANT: Lingner, Joachim
; APPLICANT: Nakamura, Toru
; APPLICANT: Chapman, Karen B.
; APPLICANT: Morin, Gregg B.
; APPLICANT: Harley, Calvin B.
; APPLICANT: Andrews, William H.
; TITLE OF INVENTION: Human Telomerase Catalytic Subunit
; NUMBER OF SEQUENCES: 727
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn*Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/974,549A
; FILING DATE: 19-NOV-1997
; CLASSIFICATION: 536
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/724,643
; FILING DATE: 01-OCT-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/844,419
; FILING DATE: 18-APR-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/846,017
; FILING DATE: 25-APR-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/851,843
; FILING DATE: 06-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/854,050
; FILING DATE: 09-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/911,312
; FILING DATE: 14-AUG-1997
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; APPLICATION NUMBER: US 08/912,951
; FILING DATE: 14-AUG-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/915,503
; FILING DATE: 14-AUG-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US97/17618
; FILING DATE: 01-OCT-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/US97/17885
; FILING DATE: 01-OCT-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Apple, Randolph Ted
; REGISTRATION NUMBER: 36,429
; REFERENCE/DOCKET NUMBER: 015389-002610US
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO.: 645:

SEQUENCE CHARACTERISTICS:
; LENGTH: 90 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA
; FEATURE:
; NAME/KEY: -
; LOCATION: 1..90
; OTHER INFORMATION: /note="oligonucleotide 2B"
; US-08-974-549A-645

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; Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:
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Align seq 1/1 to: US-08-974-549A-645 from: 1 to: 90

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seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-08-974-549A-646

seq_documentation_block:
; Sequence 646, Application US/08974549A
; Patent No. 6166178
; GENERAL INFORMATION:
; APPLICANT: Cech, Thomas R.
; APPLICANT: Lingner, Joachim
; APPLICANT: Nakamura, Toru
; APPLICANT: Chapman, Karen B.
; APPLICANT: Morin, Gregg B.
; APPLICANT: Harley, Calvin B.
; APPLICANT: Andrews, William H.
; TITLE OF INVENTION: Human Telomerase Catalytic Subunit
; NUMBER OF SEQUENCES: 727
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834

COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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; FILING DATE: 19-NOV-1997
; CLASSIFICATION: 536
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/724,643
; FILING DATE: 01-OCT-1996
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; APPLICATION NUMBER: US 08/844,419
; FILING DATE: 18-APR-1997
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; APPLICATION NUMBER: US 08/846,017
; FILING DATE: 25-APR-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/851,843
; FILING DATE: 06-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/854,050
; FILING DATE: 09-MAY-1997

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: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/911,312
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: APPLICATION NUMBER: US 08/915,503
: FILING DATE: 14-AUG-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: WO PCT/US97/17618
: FILING DATE: 01-OCT-1997
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: WO PCT/US97/17885
: FILING DATE: 01-OCT-1997
: ATTORNEY/AGENT INFORMATION:
: NAME: Apple, Randolph Ted
: REGISTRATION NUMBER: 36,429
: REFERENCE/DOCKET NUMBER: 015389-002610US
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 576-0200
: TELEFAX: (415) 576-0300
: INFORMATION FOR SEQ ID NO: 646:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 90 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: DNA
: FEATURE:
: NAME/KEY:
: LOCATION: 1..90
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: US-08-974-549A-646

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About: Results were produced by the GenCore software, version 4.5,
 Copyright (c) 1993-2000 CompuGen Ltd.

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Search information block:

Query: US-09-444-281-27
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gb.pr:AC005242	-	42.00	88.81	1.1e+04	160262	AC005242 Homo sapiens chrom
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gb.ba:AP002996 - 42.00 83.52 2.2e+04 349619 | AP002996 Mesorhizobium lo
 gb.un:M27461 - 41.00 122.64 148.45 687 | M27461 Figure 3. DNA sequenc
 gb.pr:AF172450 - 41.00 116.59 322.63 1676 | AF172450 Homo sapiens optoi
 gb.pr:AF172452 - 41.00 114.26 435.06 2363 | AF172452 Homo sapiens optoi

seq_name: gb_ph:AF109874

seq_documentation block:

LOCUS AF109874 38347 bp DNA circular PFG 15-MAR-2001
 DEFINITION Bacteriophage Tuc2009, complete genome.
 ACCSSION AF109874 L26219 L31348 L31364 L31365 L31366
 VERSION AF109874.2 GI:13346831
 KEYWORDS
 SOURCE Bacteriophage Tuc2009.
 ORGANISM Bacteriophage Tuc2009.
 Viruses; dsDNA viruses, no RNA stage; Tailed phages; Siphoviridae.
 1 (bases 24100 to 24323; 33392 to 34000; 36107 to 38347)
 Arendt,E.K., Daly,C., Fitzgerald,G.F. and van de Guchte,M.
 Molecular characterization of lactococcal bacteriophage Tuc2009 and
 identification and analysis of genes encoding lysis, a putative
 holin, and two structural proteins
 Appl. Environ. Microbiol. 60 (6), 1875-1883 (1994)

JOURNAL MEDLINE 94304164
 PUBMED 8031083
 TITLE 2 (bases 1921 to 2781)
 van de Guchte,M., Daly,C., Fitzgerald,G.F. and Arendt,E.K.
 Identification of the putative repressor-encoding gene ci of the
 temperate lactococcal bacteriophage Tuc2009

JOURNAL MEDLINE 94299176
 PUBMED 8026765
 TITLE 3 (bases 37811 to 38347; 1 to 1335)
 van de Guchte,M., Daly,C., Fitzgerald,G.F. and Arendt,E.K.
 Identification of int and attP on the genome of lactococcal
 bacteriophage Tuc2009 and their use for site-specific plasmid
 integration in the chromosome of Tuc2009-resistant Lactococcus

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 4 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Molecular analysis of the temperate lactococcal phage Tuc2009

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 5 (bases 4602 to 11597)
 McGrath,S., Seegers,J.F.M.L., Fitzgerald,G.F., van Sinderen,D. and
 van de Guchte,M.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
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JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
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JOURNAL MEDLINE 94356466
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 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
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JOURNAL MEDLINE 94356466
 PUBMED 8074513
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 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

JOURNAL MEDLINE 94356466
 PUBMED 8074513
 TITLE 6 (bases 1 to 38347)
 van Sinderen,D., van de Guchte,M., Seegers,J.F.M.L. and
 Fitzgerald,G.F.
 Direct Submission

CDS
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/db_xref="GI:5001700"
/translation="MINNVYLVGRITPDPPELRHTPONOAVGTGELAVNROFKRANGER
EADLINCIVIROQAEENAKRAKKALIGTIGRTOTRYENQOQKYYVEVAVDTROM
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alignment_scores: 46.00 Length: 9
Ratio: 5.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889
alignment_block:
US-09-444-281-27 x AF109874/rev ..
Align seg 1/1 to reverse of: AF109874 from: 1 to: 38347

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1 HISGLUAlaGluProGluAlaGluPro 9
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35925 CATAAAGCTGACAGAGCTGACCT 35899

seq_name: gb_pr:AL391821

seq_documentation_block:
LOCUS      AL391821      158408 bp      DNA      PRI      01-NOV-2000
DEFINITION Human DNA sequence from clone RP11-212822 on chromosome X, complete
sequence.
ACCESSION  AL391821
VERSION    AL391821.7
KEYWORDS   GI:11121082
SOURCE     HTG.
           human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE  1 (bases 1 to 158408)
AUTHORS   Heath, P.
TITLE     Direct Submission
JOURNAL   Submitted (31-OCT-2000) Sanger Centre, Hinxton, Cambridgeshire,
        GB10 ISA, UK. E-mail enquiries: humquerry@sanger.ac.uk Clone
        requests: clonerequests@sanger.ac.uk
        On Nov 8, 2000 this sequence version replaced gi:10944214.
        During sequence assembly data is compared from overlapping clones.
        Where differences are found these are annotated as variations
        together with a note of the overlapping clone name. Note that the
        variation annotation may not be found in the sequence submission
        corresponding to the overlapping clone, as we submit sequences with
        only a small overlap as described above.
        This sequence has been finished according to sequence map criteria
        as follows. An attempt is made to resolve all sequencing problems,
        such as compressions and repeats, but not necessarily within known
        annotated human repeat sequence elements (e.g. Alu). Where the
        sequence is ambiguous, there is an annotation using the 'unsure'
        feature key.
        The following abbreviations are used to associate primary accession
        numbers given in the feature table with their source databases:
        Em: EMBL; SW: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
        on the WORMPEP database can be found at
        http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
        was generated from part of bacterial clone contigs of human
        chromosome X, constructed by the Sanger Centre Chromosome X Mapping
        Group. Further information can be found at
        http://www.sanger.ac.uk/HGP/ChrX
        RP11-212822 is from the library RPCI-11.1 constructed at the
        Roswell Park Cancer Institute by the group of Pieter de Jong. For
        further details see http://bacpac.med.buffalo.edu/
        VECTOR: pBACE3.6
        This sequence is the entire insert of clone RP11-212822 The true
        left end of clone RP11-48719 is at 133439 in this sequence. The
        true right end of clone RP11-35618 is at 153997 in this sequence.
FEATURES             Location/Qualifiers
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                        /db_xref="taxon:9606"
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     repeat_region     295..525
                        /note="MLT1H repeat: matches 229..494 of consensus"
     repeat_region     648..677
                        /note="15 copies 2 mer ca 90% conserved"
     repeat_region     2598..2649
                        /note="26 copies 2 mer ac 90% conserved"
     repeat_region     2780..2853
                        /note="37 copies 2 mer ct 74% conserved"
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     repeat_region     4491..4655
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repeat_region 4804..5008
/note="MER58C repeat: matches 3..88 of consensus"
repeat_region 5044..5181
/note="LIM4 repeat: matches 1500..1639 of consensus"
repeat_region 5271..5414
/note="LIMC repeat: matches 2242..2385 of consensus"
repeat_region 5420..5718
/note="Alusq repeat: matches 1..298 of consensus"
repeat_region 5726..5994
/note="Alu0 repeat: matches 1..271 of consensus"
repeat_region 6011..6787
/note="LIMC repeat: matches 2356..2818 of consensus"
repeat_region 6801..7262
/note="LIPAS repeat: matches 5682..6143 of consensus"
repeat_region 7305..7750
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repeat_region 7799..8106
/note="LIM4 repeat: matches 2841..3151 of consensus"
repeat_region 8246..8761
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repeat_region 8819..9030
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repeat_region 9073..10246
/note="LIPAS repeat: matches 4979..6146 of consensus"
repeat_region 10260..10753
/note="LIM4 repeat: matches 1575..1608 of consensus"
repeat_region 11391..11509
/note="FLAM_C repeat: matches 1..132 of consensus"
repeat_region 11510..11821
/note="Alusq repeat: matches 1..312 of consensus"
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repeat_region 12408..12684
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repeat_region 12685..13112
/note="LIM4 repeat: matches 2308..3346 of consensus"
repeat_region 13191..13493
/note="LIM4 repeat: matches 3350..3653 of consensus"
repeat_region 13515..13538
/note="12 copies 2 mer at 100% conserved"
repeat_region 13539..13894
/note="LIM9 repeat: matches 4974..5334 of consensus"
repeat_region 13895..14205
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repeat_region 14206..15193
/note="LIMAS repeat: matches 5334..6305 of consensus"
repeat_region 15269..15357
/note="LIMAS repeat: matches 2644..2748 of consensus"
repeat_region 15388..15841
/note="LIP repeat: matches 4685..5138 of consensus"
repeat_region 16099..16221
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repeat_region 16280..16732
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repeat_region 17982..18300
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repeat_region 18532..18756
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repeat_region 20534..20587
/note="MIR repeat: matches 14..262 of consensus"
repeat_region 21039..21171
/note="27 copies 2 mer tt 81% conserved"
repeat_region 22602..22902
/note="Alusq/x repeat: matches 79..209 of consensus"
repeat_region 23484..23533
/note="Alusx repeat: matches 2..302 of consensus"
repeat_region 23532..23561
/note="25 copies 2 mer gt 96% conserved"
repeat_region 23877..25040
/note="10 copies 3 mer gtg 90% conserved"
repeat_region 25257..25415
/note="L2 repeat: matches 1141..2323 of consensus"
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28430. .28618
/note="MER53 repeat: matches 1. .189 of consensus"
repeat_region 28941. .29026
/note="MIR repeat: matches 112. .198 of consensus"
30445. .30809
/note="THE1B repeat: matches 1. .364 of consensus"
repeat_region 31228. .31341
/note="L2 repeat: matches 2629. .2749 of consensus"
31996. .32079
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/note="AluY repeat: matches 1. .294 of consensus"
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33673. .33723
/note="MIR repeat: matches 102. .152 of consensus"
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38390. .38732
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/note="MIR repeat: matches 8. .261 of consensus"
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42729. .42884
/note="MER5A repeat: matches 4. .184 of consensus"
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45117. .45305
/note="MER53 repeat: matches 1. .194 of consensus"
45425. .45951
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46025. .46336
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46474. .46532
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alignment_scores:
  Quality: 44.00
  Ratio: 5.500
  Percent Similarity: 88.889
  Percent Identity: 88.889

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alignment_block:
  US-09-444-281-27 x AL391821/rev

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Align seg 1/1 to reverse of: AL391821 from: 1 to: 156408

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1 HisGluAgiProGluAgiGluPro 9

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96328 CATGAGCTGAGCCAGATTGAGCCT 96302
seq_name: gb_htg:AC021098

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seq_documentation_block:
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DEFINITION Homo sapiens chromosome X clone RP11-33A2, WORKING DRAFT SEQUENCE,
18 unordered pieces.
ACCESSION AC021098.3 GI:7230836
VERSION AC021098
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 169671)
Waterston,R.H.
The sequence of Homo sapiens clone
2 (bases 1 to 169671)
Waterston,R.H.
Direct Submission
Submitted (14-JAN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Mar 13, 2000 this sequence version replaced gi:6922906.

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COMMENT

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----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
----- Project Information -----
Center project name: H.NH0033A02
----- Summary Statistics -----
Sequencing vector: M13, 81%
Sequencing vector: Plasmid, 19%
Chemistry: Dye-primer ET; 81% of reads
Chemistry: Dye-terminator Big Dye; 19% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 159126 bases at least Q40
Consensus quality: 162659 bases at least Q30
Consensus quality: 164792 bases at least Q20
Insert size: 170000; agarose-fp
Insert size: 167971; sum-of-ctrls
Quality coverage: 4.16 in Q20 bases; agarose-fp
Quality coverage: 4.21 in Q20 bases; sum-of-ctrls
-----

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

```

```

* 1 1196: contig of 1196 bp in length
* 1197 1296: gap of unknown length
* 1297 3284: contig of 1988 bp in length
* 3285 3384: gap of unknown length
* 3385 6023: contig of 2639 bp in length
* 6024 6123: gap of unknown length
* 6124 8768: contig of 2645 bp in length
* 8769 8868: gap of unknown length
* 8869 12171: contig of 3303 bp in length
* 12172 12271: gap of unknown length
* 12272 15977: contig of 3706 bp in length
* 15978 16078: gap of unknown length
* 16079 24250: contig of 8173 bp in length
* 24251 24350: gap of unknown length
* 24351 32681: contig of 8331 bp in length
* 32682 32781: gap of unknown length
* 32782 41939: contig of 9158 bp in length

```

```

* 41940 42039: gap of unknown length
* 42040 48802: contig of 6763 bp in length
* 48803 48902: gap of unknown length
* 48903 57427: contig of 8525 bp in length
* 57428 57527: gap of unknown length
* 57528 66323: contig of 8796 bp in length
* 66324 66423: gap of unknown length
* 66424 78283: contig of 11860 bp in length
* 78284 78384: gap of unknown length
* 78384 93662: contig of 15279 bp in length
* 93663 93762: gap of unknown length
* 93763 111526: contig of 17764 bp in length
* 111527 129714: contig of 18088 bp in length
* 129715 129814: gap of unknown length
* 129815 144537: contig of 14723 bp in length
* 144538 146571: gap of unknown length
* 144638 169671: contig of 25034 bp in length.
location/Qualifiers

```

FEATURES

source

```

1. 169671
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-33A2"
1. 1196
/note="assembly_name:Contig7"
misc_feature
1297..3284
/note="assembly_name:Contig8"
misc_feature
3385..6023
/note="assembly_name:Contig9"
misc_feature
6124..8768
/note="assembly_name:Contig10"
misc_feature
8869..12171
/note="assembly_name:Contig11"
misc_feature
12272..15977
/note="assembly_name:Contig12"
misc_feature
16078..24250
/note="assembly_name:Contig13"
misc_feature
24351..32681
/note="assembly_name:Contig14"
misc_feature
32782..41939
/note="assembly_name:Contig15"
misc_feature
42040..48802
/note="assembly_name:Contig16"
misc_feature
clone_end:Sp6
vector_side:right"
48903..57427
/note="assembly_name:Contig17"
misc_feature
clone_end:T7
vector_side:right"
57528..66323
/note="assembly_name:Contig18"
misc_feature
66424..78283
/note="assembly_name:Contig19"
misc_feature
78384..93662
/note="assembly_name:Contig20"
misc_feature
93763..111526
/note="assembly_name:Contig21"
misc_feature
111627..129714
/note="assembly_name:Contig22"
misc_feature
129815..144537
/note="assembly_name:Contig23"
misc_feature
144638..169671
/note="assembly_name:Contig24"
misc_feature
BASE COUNT 53919 a 31234 c 31915 g 50898 t 1705 others
ORIGIN

```

alignment_scores: 44.00 Length: 9
 Ratio: 5.500 Gaps: 0
 Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-09-444-281-27 x AC021098 ..

Align seg 1/1 to: AC021098 from: 1 to: 169671

```

1 HiscLualagiupProglualagiPro 9
|||||
44084 CATGAGCTGAGCCAGATTGACCT 44110

```

seq_name: gb_ro:RRXPFPRT

seq_documentation_block:

LOCUS RRPXPFPRT 966 bp mRNA ROD 01-OCT-1999
 DEFINITION R. rattus mRNA for Pxf protein.
 ACCESSION Y09049
 VERSION Y09049.1 GI:6010290
 KEYWORDS Pxf protein.
 SOURCE black rat.
 ORGANISM Rattus rattus
 Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.
 1 (bases 1 to 966)
 AUTHORS Kammerer, S.
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 966)
 AUTHORS Kammerer, S.
 TITLE Direct Submission
 JOURNAL Submitted (22-OCT-1996) S. Kammerer, Kinderkrankenhaus, Labor f.
 Molekulare Biologie, Lindwurmstr. 4, D-80337 Muenchen, FRG
 Location/Qualifiers

FEATURES

source

```

1. 966
/organism="Rattus rattus"
/db_xref="taxon:10117"
/cell_type="myocardocyte"
<1..75
/number=1
join(6..75,76..185,186..351,352..437,438..599,600..776,
777..821,822..905)
/number=2
/number=3
/number=4
/number=5
/number=6
/number=7
/number=8

```

BASE COUNT 257 a 247 c 270 g 192 t
ORIGIN

alignment_scores:

Quality: 43.00 Length: 11
Ratio: 3.909 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 63.636

alignment_block:

US-09-444-281-27 x RRPXPPT

Align seg 1/1 to: RRPXPPT from: 1 to: 966

1 HisGluAaIupProGluAaIupProIleMet 11
|||||
524 CATCATGACAGACCTCATGACAGACCTGTGTC 556

seq_name: gb_da:AF127222

seq_documentation_block:

LOCUS AF127222 2501 bp DNA BCI 07-FEB-2000
DEFINITION Pseudomonas fluorescens, outer membrane heme receptor Pfhr (pfhr)
gene, complete cds.
ACCESSION AF127222
VERSION AF127222.1 GI:4838476

KEYWORDS

Pseudomonas fluorescens.
Pseudomonas fluorescens.
Bacteria: Proteobacteria; gamma subdivision: Pseudomonadaceae;

ORGANISM

Pseudomonas.

1 (bases 1 to 2501)
Ochsner, U.A., Johnson, Z., and Vasil, M.L.
Genetics and regulation of two distinct haem-uptake systems, phu
and has, in Pseudomonas aeruginosa
Microbiology 146 (Pt 1), 185-198 (2000)

20121752

10658665

2 (bases 1 to 2501)

Ochsner, U.A. and Vasil, M.L.
Direct Submission
Submitted (10-FEB-1999) Microbiology, University of Colorado Health
Sci Ctr, 4200 E. Ninth Avenue, Denver, CO 80262, USA

FEATURES

Source

1..2501
/organism="Pseudomonas fluorescens"
/strain="ATCC 15453"
/db_xref="taxon:294"
/db_xref="ATCC:15453"
195..2465
/gene="pfhr"
195..2465
/gene="pfhr"
/function="heme uptake"
/codon_start=1
/transl_table=1
/product="outer membrane heme receptor pfhr"
/protein_id="AAD31012.1"
/db_xref="GI:4838477"
/translation="MGTLSTKLTNTNIVLEKRSAGSALALQOVTSATREEDVNS
VPSVSVDRALDELROHANNIRELVEPVSGAGRSNAGNINIGIDVDVLD
VDGVEVDFNFGPAKTRNRYVDEIYKREIIRGASALYSSAIGAGVSTTLD
DDIIRKPDGVCARLKTGSSADESMITSGTFRAGVODEDLHLHSQRNHTESDGN
NATGIARAGAPEDARTIVLAKLGMNIGDNRILGITEKFKODVDNLKNAVGPFTG
GRGMILYDRGNPTITREPGLENTALESPADRIKTSLNLYIATDQTTAIIYA
GRVLRFDLTYEEKOWFADLDKASISLGETDQVYGTLLKQKATGSEGSASCL
AIGACATAGSPASDSVSKASDPPTINTISLFAODITMDKTEFLDVAARYDT
RLKRLIOEFLNTVPGATVVDGDKTWNVTPKFGITVALITNNTYEGQYACGFRP
PSAKIYGRFNILNGYVNEPDLKRETSKIGTIGRKDESGFDIAYYKTRDP
IDEXRPAAGTVEOPAVANIKRAIKGVKAGRLNLDLGAPEKLYQGSVAAYYKRN
DDNEPLNSVPLGVGLGDQNGGLVSMVTVKQNRVDSITTFHAPDGTGDPFR
TPGGIIDLVAIVKSVKDVYVNGGLYMLTDKKWNMDVRSYDSVAGVGPANLDR
LTQGRNFAINVIDMI"

BASE COUNT 561 a 874 c 698 g 368 t
ORIGIN

alignment_scores:

Quality: 43.00 Length: 10
Ratio: 4.778 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-09-444-281-27 x AF127222/rev

Align seg 1/1 to reverse of: AF127222 from: 1 to: 2501

1 HisGluAaIupProGluAaIupProIle 10
|||||
1366 CACGAGCGCAGCCTTCGCTGACGCGGTG 1337

seq_name: gb_htg:AC015790

seq_documentation_block:

LOCUS AC015790 79795 bp DNA HTG 13-JUL-2000
DEFINITION Homo sapiens clone RP11-2L12, LOW-PASS SEQUENCE SAMPLING.
ACCESSION AC015790
VERSION AC015790.2 GI:9109862
KEYWORDS HTG; HTGS_PHASE0.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

1 (bases 1 to 79795)

Birren, B., Linton, L., Nusbaum, C. and Lander, E.
Homo sapiens, clone RP11-2L12
Unpublished

2 (bases 1 to 79795)

Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, M.,
Baldwin, D., Barna, N., Beckery, R., Boguslavsky, L., Boukhalter, B.,
Brown, A., Castle, A., Colangelo, M., Collins, S., Collymore, A.,
Cooke, P., DeArrellano, K., Dewar, K., Domino, M., Donelan, L., Doyle, M.,
Ferreira, P., FitzHugh, W., Forrest, C., Funke, R., Gage, D.,
Galagan, J., Gardyna, S., Grant, G., Hagos, B., Hearford, A., Horton, L.,
Howland, J.C., Johnson, R., Jones, C., Kann, L., Kartas, A., Klein, J.,
Lencock, J., Liu, C., Locke, K., Macdonald, P., Marquis, N.,
McEwan, P., McGuck, A., McKernan, K., McLaughlin, J., Meldrum, J.,
Morrison, J., Naylor, J., Norman, C.H., O'Connor, T., O'Donnell, P.,
Peterson, K., Pollara, V., Riley, R., Roy, A., Santos, R., Severy, P.,
Stange-Thomann, N., Stojanovic, N., Subramanian, A., Talamas, J.,
Testaye, S., Tirrell, A., Vassiliev, H., Vo, A., Wheeler, J., Wu, X.,
Wyman, D., Ye, W.J., Zimmer, A. and Zody, M.

Direct Submission
Submitted (17-NOV-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 13, 2000 this sequence version replaced gi:6446897.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: L1352
Center clone name: 2.L.12

NOTE: This record contains 80 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone

* Will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
1
* 902 1001: gap of 100 bp in length
* 1002 1910: contig of 909 bp in length
* 1911 2010: gap of 100 bp
* 2011 2938: contig of 928 bp in length
* 2939 3038: gap of 100 bp
* 3039 3033: contig of 895 bp in length
* 3034 4033: gap of 100 bp
* 4034 4931: contig of 898 bp in length
* 4932 5031: gap of 100 bp
* 5032 5905: contig of 874 bp in length
* 5906 6005: gap of 100 bp
* 6006 6884: contig of 889 bp in length
* 6895 6994: gap of 100 bp
* 6995 7902: contig of 908 bp in length
* 7903 8002: gap of 100 bp
* 8003 8887: contig of 885 bp in length
* 8888 9879: gap of 100 bp
* 9880 9979: gap of 100 bp
* 9980 10888: contig of 909 bp in length
* 10889 10988: gap of 100 bp
* 10989 11908: contig of 920 bp in length
* 11909 12008: gap of 100 bp
* 12009 12888: contig of 880 bp in length
* 12889 12988: gap of 100 bp
* 12989 13882: contig of 894 bp in length
* 13883 13982: gap of 100 bp
* 13883 14889: contig of 907 bp in length
* 14890 14989: gap of 100 bp
* 14990 15897: contig of 908 bp in length
* 15898 15997: gap of 100 bp
* 15998 16890: contig of 893 bp in length
* 16891 16990: gap of 100 bp
* 16991 17891: contig of 901 bp in length
* 17892 17991: gap of 100 bp
* 17992 18891: contig of 900 bp in length
* 18892 18991: gap of 100 bp
* 18992 19894: contig of 903 bp in length
* 19895 19994: gap of 100 bp
* 19995 20860: contig of 866 bp in length
* 20861 20960: gap of 100 bp
* 20961 21867: contig of 907 bp in length
* 21868 21967: gap of 100 bp
* 21868 22868: contig of 901 bp in length
* 22869 22968: gap of 100 bp
* 22969 23859: contig of 891 bp in length
* 23860 23959: gap of 100 bp
* 23960 24832: contig of 873 bp in length
* 24833 24932: gap of 100 bp
* 24933 25833: contig of 901 bp in length
* 25834 25933: gap of 100 bp
* 25934 26845: contig of 912 bp in length
* 26846 26945: gap of 100 bp
* 26946 27854: contig of 909 bp in length
* 27855 27954: gap of 100 bp
* 27955 28864: contig of 910 bp in length
* 28865 28964: gap of 100 bp
* 28965 29856: contig of 892 bp in length
* 29857 29956: gap of 100 bp
* 29957 30835: contig of 879 bp in length
* 30836 30935: gap of 100 bp
* 30936 31843: contig of 908 bp in length
* 31844 31943: gap of 100 bp
* 31944 32864: contig of 921 bp in length
* 32865 32964: gap of 100 bp
* 32965 33849: contig of 885 bp in length
* 33850 33949: gap of 100 bp
* 33950 34823: contig of 874 bp in length
* 34824 34923: gap of 100 bp

* 34924 35815: contig of 892 bp in length
* 35816 35915: gap of 100 bp
* 35916 36826: contig of 911 bp in length
* 36827 36926: gap of 100 bp
* 36927 37836: contig of 910 bp in length
* 37837 37936: gap of 100 bp
* 37937 38837: contig of 901 bp in length
* 38838 38937: gap of 100 bp
* 38938 39818: contig of 881 bp in length
* 39819 39918: gap of 100 bp
* 39919 40829: contig of 911 bp in length
* 40830 40929: gap of 100 bp
* 40930 41819: contig of 890 bp in length
* 41820 41919: gap of 100 bp
* 41920 42789: contig of 870 bp in length
* 42790 42889: gap of 100 bp
* 42890 43767: contig of 878 bp in length
* 43768 43867: gap of 100 bp
* 43868 44753: contig of 886 bp in length
* 44754 44853: gap of 100 bp
* 44854 45777: contig of 924 bp in length
* 45778 45877: gap of 100 bp
* 45878 46780: contig of 903 bp in length
* 46781 46880: gap of 100 bp
* 46881 47802: contig of 922 bp in length
* 47803 47902: gap of 100 bp
* 47803 48803: contig of 901 bp in length
* 48804 48903: gap of 100 bp
* 48904 49807: contig of 904 bp in length
* 49808 49907: gap of 100 bp
* 49908 50790: contig of 883 bp in length
* 50791 50890: gap of 100 bp
* 50891 51810: contig of 920 bp in length
* 51811 51910: gap of 100 bp
* 51911 52812: contig of 902 bp in length
* 52813 52912: gap of 100 bp
* 52913 53796: contig of 884 bp in length
* 53797 53896: gap of 100 bp
* 53897 54818: contig of 922 bp in length
* 54819 54918: gap of 100 bp
* 54919 55819: contig of 901 bp in length
* 55820 55919: gap of 100 bp
* 55920 56836: contig of 917 bp in length
* 56837 56936: gap of 100 bp
* 56937 57850: contig of 914 bp in length
* 57851 57950: gap of 100 bp
* 57951 58874: contig of 924 bp in length
* 58875 58974: gap of 100 bp
* 58975 59847: contig of 873 bp in length
* 59848 59947: gap of 100 bp
* 59948 60838: contig of 891 bp in length
* 60839 60938: gap of 100 bp
* 60939 61829: contig of 891 bp in length
* 61830 61929: gap of 100 bp
* 61930 62806: contig of 877 bp in length
* 62807 62906: gap of 100 bp
* 62907 63784: contig of 878 bp in length
* 63785 63884: gap of 100 bp
* 63885 64792: contig of 908 bp in length
* 64793 64892: gap of 100 bp
* 64893 65789: contig of 897 bp in length
* 65790 65889: gap of 100 bp
* 65890 66771: contig of 882 bp in length
* 66772 66871: gap of 100 bp
* 66872 67778: contig of 907 bp in length
* 67779 67878: gap of 100 bp
* 67879 68780: contig of 902 bp in length
* 68781 68880: gap of 100 bp
* 68881 69764: contig of 884 bp in length
* 69765 69864: gap of 100 bp
* 69865 70743: contig of 879 bp in length
* 70744 70843: gap of 100 bp
* 70844 71756: contig of 913 bp in length

* 71757 71856: gap of 100 bp
* 71857 72761: contig of 905 bp in length

alignment_scores:
Quality: 43.00 Length: 10
Ratio: 4.300 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 80.000

Alignment block:

US-09-444-281-27 x AC015790/rev

Align seg 1/1 to reverse of: AC015790 from: 1 to: 79795

2 GUAAGAGUProGUAGAGUProIleket 11
||||:|||||
4914 GAAGCTCCGACGACGACCATCATG 4885

seq_name: gb.pr:AL138709

seq_documentation_block:

LOCUS AL138709 96874 bp DNA PRI 08-AUG-2001
DEFINITION Human DNA sequence from clone RP11-460G11 on chromosome 13,
complete sequence.

ACCESSION AL138709

VERSION AL138709.19 GI:15141998

KEYWORDS HTG.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

AUTHORS Laid,C.

TITLE Direct Submission

JOURNAL Submitted (08-AUG-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk

COMMENT

On Aug 9, 2001 this sequence version replaced gi:14626037.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats: all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em: EMBL; SW:
SWISSPROT; Tr: TrEMBL; Wp: WormPEP; Information on the WormPEP
database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
Chromosome 13, constructed by the Sanger Centre Chromosome 13
Mapping Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr13>
RP11-460G11 is from the library RP11-11.2 constructed by the group
of Pieter de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>

VECTOR: pACe3.6
IMPORTANT: This sequence is not the entire insert of clone
RP11-460G11. It may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true right end of clone RP11-460G11 is at 96874 in this
sequence. The true right end of clone RP11-335618 is at 2000 in
this sequence.

FEATURES

Source

Location/Qualifiers
1..96874
/organism="Homo sapiens"
/db_xref="taxon:9606"

/chromosome="13"
/clone_id="RP11-460G11"
/clone_lib="RP11-11.2"
638..775
/note="AlusB/q repeat: matches 159..296 of consensus"
2734..2953
/note="MIR repeat: matches 20..252 of consensus"
3412..3570
/note="Charlied4 repeat: matches 1740..1907 of consensus"
4223..4292
/note="MIR repeat: matches 133..205 of consensus"
4354..4475
/note="MER5B repeat: matches 1..129 of consensus"
4980..5129
/note="MER5A repeat: matches 13..164 of consensus"
5146..5284
/note="MIR repeat: matches 9..148 of consensus"
5314..5483
/note="Charlied4 repeat: matches 17..174 of consensus"
5484..5788
/note="AluY repeat: matches 5..309 of consensus"
5789..6029
/note="Charlied4 repeat: matches 174..457 of consensus"
6536..6602
/note="L2 repeat: matches 2675..2743 of consensus"
6719..7125
/note="MLR2FB repeat: matches 4..414 of consensus"
7364..7568
/note="MIR repeat: matches 37..251 of consensus"
10015..10064
/note="MIR repeat: matches 41..90 of consensus"
10027..10085
/note="MER3 repeat: matches 144..204 of consensus"
10086..10325
/note="L1M1 repeat: matches 5550..5792 of consensus"
10326..10634
/note="AlusX repeat: matches 1..309 of consensus"
10635..11160
/note="L1M1 repeat: matches 5792..6304 of consensus"
11161..11179
/note="MER3 repeat: matches 130..144 of consensus"
11182..11347
/note="MER3 repeat: matches 1..167 of consensus"
11713..11780
/note="4 copies 17 mer 86% conserved"
11787..11881
/note="L2 repeat: matches 2575..2699 of consensus"
12183..12292
/note="MIR repeat: matches 126..246 of consensus"
13008..13053
/note="23 copies 2 mer tc 73% conserved"
13063..13218
/note="6 copies 26 mer 67% conserved"
13065..13178
/note="57 copies 2 mer tc 67% conserved"
13674..13759
/note="MIR repeat: matches 78..165 of consensus"
15493..15544
/note="26 copies 2 mer ac 90% conserved"
16508..17150
/note="MER41B repeat: matches 3..635 of consensus"
18228..18286
/note="tRNA-Ile-ATC repeat: matches 14..74 of consensus"
18765..19215
/note="L2 repeat: matches 2290..2748 of consensus"
19216..19985
/note="LTR39 repeat: matches 1..794 of consensus"
19986..20070
/note="L2 repeat: matches 2208..2290 of consensus"
21039..21339
/note="Alusq repeat: matches 1..301 of consensus"
21350..21658
/note="Alusq repeat: matches 1..307 of consensus"


```

repeat_region 21674..21731
/note="MER53 repeat: matches 127..186 of consensus"
repeat_region 21732..22043
/note="AluSx repeat: matches 1..312 of consensus"
repeat_region 22044..22175
/note="MER53 repeat: matches 6..127 of consensus"
repeat_region 22862..23024
/note="MIR repeat: matches 87..248 of consensus"
repeat_region 24844..25143
/note="AluSx repeat: matches 1..302 of consensus"
repeat_region 26057..26200
/note="MER5A repeat: matches 17..187 of consensus"
repeat_region 27843..28278
/note="MLTID repeat: matches 1..466 of consensus"
repeat_region 28375..28436
/note="L2 repeat: matches 2645..2710 of consensus"
repeat_region 30397..30480
/note="2 copies 42 mer 90% conserved"
repeat_region 31856..32054
/note="MIR repeat: matches 37..261 of consensus"
repeat_region 32358..32528
/note="L2 repeat: matches 2092..2279 of consensus"
repeat_region 32608..32648
/note="MLTID repeat: matches 201..240 of consensus"
repeat_region 32649..32947
/note="AluJo repeat: matches 1..294 of consensus"
repeat_region 32948..33144
/note="MLTID repeat: matches 1..201 of consensus"
repeat_region 33183..33230
/note="L2 repeat: matches 2677..2724 of consensus"
repeat_region 34446..34647
/note="MER2 repeat: matches 1..211 of consensus"
repeat_region 34678..34757
/note="MER2 repeat: matches 264..344 of consensus"
repeat_region 38187..38646
/note="MLTIC repeat: matches 6..466 of consensus"
repeat_region 38848..38958
/note="MER81 repeat: matches 3..114 of consensus"
repeat_region 40042..40152
/note="L2 repeat: matches 2381..2492 of consensus"
repeat_region 41116..41251
/note="MIR repeat: matches 15..151 of consensus"
repeat_region 42941..43367
/note="LTP16B repeat: matches 39..464 of consensus"
repeat_region 44219..44302
/note="MLTID repeat: matches 242..318 of consensus"
repeat_region 46089..47392
/note="LTPA2 repeat: matches 3642..4945 of consensus"
repeat_region 47393..48587
/note="LTPA2 repeat: matches 4952..6146 of consensus"
repeat_region 48647..48781
/note="MIR repeat: matches 85..234 of consensus"
repeat_region 49096..49396
/note="AluSg repeat: matches 1..296 of consensus"
repeat_region 49623..51400
/note="LTPA7 repeat: matches 4358..6143 of consensus"
repeat_region 52702..53151
/note="MER39 repeat: matches 13..455 of consensus"
repeat_region 53233..53451
/note="MER72 repeat: matches 445..668 of consensus"
repeat_region 53452..53750
/note="AluSg repeat: matches 1..294 of consensus"
repeat_region 53751..54215
/note="MER72 repeat: matches 1..445 of consensus"
repeat_region 54223..54364
/note="L2 repeat: matches 2563..2750 of consensus"
repeat_region 54866..55171
/note="AluY repeat: matches 2..303 of consensus"
repeat_region 55239..55292
/note="27 copies 2 mer tt 70% conserved"
repeat_region 57830..58112
/note="AluSg repeat: matches 3..304 of consensus"
repeat_region 58224..58516

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alignment_scores:
  Quality: 43.00      Length: 11
  Ratio: 4.778      Gaps: 0
  Percent Similarity: 81.818      Percent Identity: 63.636
alignment_block:
  US-09-444-281-27 x AL138709/rev ..
Align seg 1/1 to reverse of: AL138709 from: 1 to: 96874
1 HtSGuAlagluProgluAlagluProtleMet 11
||||| ||||||| |||||||:
66249 CACGAGCAGACCTGACCCGAGCCCTGGTT 66217
seq_name: gb_hlg:AC019290
seq_documentation_block:
LOCUS      AC019290      176712 bp      DNA
DEFINITION Homo sapiens clone RP11-649M10, WORKING DRAFT SEQUENCE, 15
unordered pieces.
ACCESSION      AC019290
VERSION      AC019290.4 GI:10045469
KEYWORDS      HTG: HTGS_PHASE1; HTGS_DRAFT.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 176712)
REFERENCE      Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE      Homo sapiens, clone RP11-649M10
JOURNAL      unpublished
REFERENCE      2 (bases 1 to 176712)
AUTHORS      Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barua,N., Beckerly,R., Beda,F.,
Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,
Choeplel,Y., Collangelo,M., Collins,S., Collymore,A., Cooke,P.,
Dearellano,K., Dewar,K., Domino,M., Doyle,M., Feneator,J.,
Ferreira,P., Fitzhugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Landers,T., Lehoczeky,J., Levine,R., Lieu,C., Liu,G., Locke,K.,
Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Morrow,J., Naylor,J.,
Norman,C.H., O'Connor,T., O'Donnell,P., Oliver,T.M., Peterson,K.,
Pierre,N., Pisanl,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (31-DEC-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA.
On Sep 9, 2000 this sequence version replaced gi:6721321.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: MIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L4312
Center clone name: 649_M_10
----- Summary Statistics
Sequencing vector: M13: M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 167161 bases at least Q40
Consensus quality: 171786 bases at least Q30
Consensus quality: 173554 bases at least Q20

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Insert size: 182000; agarose-fp
 Insert size: 175312; sum-of-contigs
 Quality coverage: 4.3 in Q20 bases; agarose-fp
 Quality coverage: 4.5 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 15 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 31590: contig of 31590 bp in length
 31591 31690: gap of 100 bp
 31691 33087: contig of 1397 bp in length
 33088 33187: gap of 100 bp
 33188 34887: contig of 1700 bp in length
 34888 34987: gap of 100 bp
 34988 37644: contig of 2657 bp in length
 37645 37744: gap of 100 bp
 37745 40979: contig of 3235 bp in length
 40980 41079: gap of 100 bp
 41080 46341: contig of 5262 bp in length
 46342 46441: gap of 100 bp
 46442 53992: contig of 7551 bp in length
 53993 54092: gap of 100 bp
 54093 60116: contig of 6024 bp in length
 60117 60216: gap of 100 bp
 60217 68183: contig of 7967 bp in length
 68184 68283: gap of 100 bp
 68284 76928: contig of 8645 bp in length
 76929 77028: gap of 100 bp
 77029 95627: contig of 18599 bp in length
 95628 95727: gap of 100 bp
 95728 115794: contig of 20067 bp in length
 115795 115894: gap of 100 bp
 115895 13115: contig of 27221 bp in length
 143116 143215: gap of 100 bp
 143216 174379: contig of 31164 bp in length
 174380 174479: gap of 100 bp
 174480 176712: contig of 2233 bp in length.

FEATURES

Source
 1. 176712
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="RP11-649M10"
 /clone_lib="RPC1-11 Human Male BAC"
 1. 31590
 /note="assembly-fragment"
 clone_end:SP6
 vector_side:left
 31691. 33087
 /note="assembly-fragment"
 33188. 34887
 /note="assembly-fragment"
 34988. 37644
 /note="assembly-fragment"
 37745. 40979
 /note="assembly-fragment"
 41080. 46341
 /note="assembly-fragment"
 46442. 53992
 /note="assembly-fragment"
 54093. 60116
 /note="assembly-fragment"
 60217. 68183
 /note="assembly-fragment"
 68284. 76928
 /note="assembly-fragment"
 77029. 95627
 /note="assembly-fragment"
 95728. 115794

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 115895..143115
 /note="assembly-fragment"
 misc_feature 143216..174379
 /note="assembly-fragment"
 misc_feature 174480..176712
 /note="assembly-fragment"
 clone_end:17
 vector_side:right

BASE COUNT 53093 a 35421 c 35789 g 51008 t 1401 others
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alignment_scores:
 Quality: 43.00 Length: 11
 Ratio: 4.300 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 72.727

alignment_block:
 US-09-444-281-27 x AC019290/rev ..

Align seq 1/1 to reverse of: AC019290 from: 1 to: 176712

1 HisgluAlaGluProGluAlaGluProIleMet 11
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 65329 CATGAGGCTCCGCCAGACGATGCCATCATG 65297

seq_name: gb_htg:AC016270

seq_documentation_block:
 LOCUS AC016270 189022 bp DNA

DEFINITION Homo sapiens clone RP11-17111, WORKING DRAFT SEQUENCE, 15 unordered pieces

AC016270

VERSION AC016270.4 GI:7248975

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 189022)

Bliren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens, clone RP11-17111

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

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TITLE

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

JOURNAL

TITLE

REFERENCE

AUTHORS

Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIRB
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 Project Information
 Center project name: L3645

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Center clone name: 17.1.11
----- Summary Statistics
Sequencing vector: M13; M77815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 182484 bases at least Q40
Consensus quality: 184881 bases at least Q40
Consensus quality: 186037 bases at least Q20
Insert size: 186000; agarose-1p
Insert size: 187622; sum-of-contigs
Quality coverage: 5.8 in Q20 bases; agarose-1p
Quality coverage: 5.8 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 15 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 1585: contig of 1585 bp in length
* 1586 1685: gap of 100 bp
* 1586 3061: contig of 1376 bp in length
* 3062 3161: gap of 100 bp
* 3162 7504: contig of 4343 bp in length
* 7505 7604: gap of 100 bp
* 7605 10536: contig of 2932 bp in length
* 10537 10636: gap of 100 bp
* 10637 13824: contig of 3188 bp in length
* 13825 13924: gap of 100 bp
* 13925 18777: contig of 4853 bp in length
* 18778 18877: gap of 100 bp
* 18878 29087: contig of 10210 bp in length
* 29088 29187: gap of 100 bp
* 29188 40374: contig of 11187 bp in length
* 40375 40474: gap of 100 bp
* 40475 55299: contig of 14825 bp in length
* 55300 55399: gap of 100 bp
* 55400 67577: contig of 12178 bp in length
* 67578 67677: gap of 100 bp
* 67678 86116: contig of 18439 bp in length
* 86117 86216: gap of 100 bp
* 86217 104553: contig of 18337 bp in length
* 104554 104653: gap of 100 bp
* 104654 125313: contig of 20660 bp in length
* 125314 125413: gap of 100 bp
* 125414 154037: contig of 28624 bp in length
* 154038 154137: gap of 100 bp
* 154138 189022: contig of 34885 bp in length.

FEATURES
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/clone_RP11-17111"
/clone_11b="RPCT-11 Human Male BAC"
1..1585
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1686..3061
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3162..7504
/note="assembly-fragment"
7605..10536
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10637..13824
/note="assembly-fragment"
13925..18777
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18878..29087
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29188..40374
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clone_end:5p6

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clone_end:77
vector_side:right"
125414..154037
/note="assembly_fragment"
154138..189022
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BASE COUNT          55798 a 38604 c 38373 g 54845 t 1402 others
ORIGIN
alignment_scores:
Quality:          43.00          Length:          11
Ratio:            4.778          Gaps:            0
Percent Similarity: 81.818          Percent Identity: 63.636
alignment_block:
US-09-444-281-27 x AC016270 ..
Align seg 1/1 to: AC016270 from: 1 to: 189022
1 HisGluAlaGluProGluAlaGluProIleMet 11
||||| ||||||| |||||||:
43795 CACGACCGAGCGCTGAGCCGAGCCCGTGT 43827
seq_name: gb_in:AMAPID22
seq_documentation_block:
LOCUS          607 bp          INV          03-MAY-1993
DEFINITION          A.mellifera Apid22 mRNA.
ACCESSION          X72576
VERSION          X72576.1 GI:297064
KEYWORDS          antibiotic peptide; apid22 gene; Apidaein; precursor protein.
SOURCE          honeybee.
ORGANISM          Apis mellifera
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita;
Aculeata; Apoidea; Apidae; Apis.
1 (bases 1 to 607)
Casteels-Josson, K.
Direct Submission
Submitted (04-MAR-1993) K. Casteels-Josson, Memorial Sloan
Kettering Cancer Center, 1275 York Avenue, New York 10021, NY, USA
2 (bases 1 to 607)
Casteels-Josson, K., Capaci, T., Casteels, P. and Tempst, P.
Apidaein multipetide precursor structure: a putative mechanism
for amplification of the insect antibacterial response
EMBO J. 12 (4), 1569-1578 (1993)
932323697
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location/Qualifiers
1..607
/organism="Apis mellifera"
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/gene="ITGAF"

join=573...778, Y12383.1:179...206, Y12383.1:307...386, Y12384.1:87...342, Y12385.1:14...145, Y12385.1:391...510, Y12386.1:113...320, Y12386.1:1423...616, Y12386.1:859...947, Y12386.1:1057...1184, Y12386.1:1311...1406, Y12386.1:11547...1608, Y12388.1:1710...1879, Y12387.1:110...259, Y12388.1:1110...225, Y12388.1:1622...814, Y12389.1:109...189, Y12389.1:1276...355, Y12389.1:1454...528, Y12389.1:667...769, Y12389.1:1256...1432, Y12389.1:1597...1725, Y12390.1:64...177, Y12390.1:381...479, Y12390.1:1567...692, Y12392.1:153...283)

/gene="ITGAF"

/function="Tiamin receptor"

/note="B variant"

/codon_start=1

/label=ITGCD5B

/product="Integrin alpha 7"

/protein_id="CAA73023.1"

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/translation="MARIPRCDFELRPFGIYYITSLSLAGLEPLPAIAEINLDVMAIRK BCEPSDFEFVSALHROLOPRQSWLVGAPALALPGQARNTGGLFACPLSEETD CYRVIDIRCAVNOJESKENOMLVGVSRSQPGCKIVTCAHRESQRDOALETRDVI GTCFVLSODLAIIRDELDEGEMKFCGRPGDHOFPCOQGAATFSPSHYVFCAGP TYNMKGTAVELCAAGSPDLAHLDDGPEAAGEKRODRLPIVPAANSYGLGLFYINID SSDPOLVITKRSITLIDPADRLTGMGDLTLANSYGFSDSGKGLMRSELSFVGAAPRANH KCAVYILRKDSASRLIPEVYLSGERLTSGFSGSLAVITDLNDGADDLVGAFFERO BELUGAVVYINQMGGHMDISPLRTICGSDSMFGLISLVLGDLNDGDPDLVAGAPD DGKGVETIYHGSSLVGVAKRSQVLEGEAVGICKFSGVLSGLDNDNHNHVDLVGSLAD TAALEFRRARVILHVSQEIFDPRAIDLEPNCMDGRLLVCVDICTCSYVAPSSYSV ALNDVLDGVTGDERLRGQVRYVTELRGDLDELHOSSGYWMLKHODRVCQDTFOLE NKMDLRAIVITLISYGLRPRPLGRQAPRGELPTVAPILNAHPSORREIHFRLKCGC ODKTCOSNLQELRYOFCRSRISDTEFOALPMDLDGTRALFALSOGPFIETLVNLP S DRSQADDDDDHEAQLVTLTPASLRTSGVRLDSVEKPLVCLNSASHVECELGPNM KRCAGVETPLISTSGITETIETPELKKLALATISOEDLPVLSRAHVEITELSTGCV ATPOOLFEGYKGEKESAMRESRDVSGSYEYVATVNOCSLTMLISAFINIMWPEIA NGKMLTLPKRVLEEGSGGGRKICSPRNILQLDVDRRRELQSGOPEOPEPEKV EESTSWMPVSAEKRNVLDDCACTAKVCVECPILXSPDRAVLHVMGTLNSTLEE YMAVSLSEYVANTVNTVSSIKNLRLDSTVIPMWYILDPAAVVEGPMWYILAV LAGLIVLEAVTLILWLKLGFEFKRAKHPREAVPYAHVKIIPREDROOFKKEKGTIOKRN WNSQMGSDAPRILIAADWHPDELGPDHVPVATA"

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/gene="ITGAF"

/function="Tiamin receptor"

/note="A variant"

/codon_start=1

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/db_xref="SPTRMBL:088731"

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sig-peptide

CDS

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ORIGIN	SRR199PCPCT10			
alignment_scores:				
Quality:	42.00	Length:	10	
Ratio:	4.667	Gaps:	0	
Percent Similarity:	90.000	Percent Identity:	70.000	
alignment_block:				
US-09-444-281-27 x MMV12380	..			
Align seg 1/1	to: MMV12380	from: 1	to: 915	
2	GUUAGluProGluuAGluProIleMet	11		
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seq_documentation_block:				
LOCUS	AMAPID73	1010 bp	mRNA	INV
DEFINITION	Apis mellifera Apid73 mRNA.			
ACCESSION	X72577			
VERSION	X72577.1 GI:297066			
KEYWORDS	antibiotic peptide; apid73 gene; Apidaecin; precursor protein.			
SOURCE	honeybee.			
ORGANISM	Apis mellifera			
REFERENCE	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Hymenoptera; Apocrita; Aculeata; Apoidea; Apidae; Apis.			
AUTHORS	Casteels-Josson, K., Capaci, T., Casteels, P. and Tempst, P.			
TITLE	Apidaecin multipptide precursor structure: a putative mechanism for amplification of the insect antibacterial response			
JOURNAL	EMBO J. 12 (4), 1569-1578 (1993)			
MEDLINE	93232697			
REFERENCE	2 (bases 1 to 1010)			
AUTHORS	Casteels-Josson, K.			
TITLE	Direct Submission			
JOURNAL	Submitted (04-MAR-1993) K. Casteels-Josson, Memorial Sloan			
FEATURES	Kettering Cancer Center, 1275 York Avenue, New York 10021, NY, USA			
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	/db_xref="GI:4539289"			
	/db_xref="SWISS-PROT:Q06602"			
	/translation="KFNALALVVTVAFAVGNTNLDPPTRPRRLREAPKPEALPEGNNN			
	RPVIVIPRPPHPRRLREAPPEALPEGNNAVRVYIQPPRPPHRLREDELPEPNNRN			

VTISPPRPRLRREAEPEAEPCGNNRPVYIPQRPRLRREAELEAEPCGNNRPVY
ISQRPRLRREAEPEAEPCGNNRPVYIPQRPRLRREAELEAEPCGNNRPVYIP
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66. .863
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misc_signal
866. .870
/note="ATTTA"
902. .906
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906. .911
polya_signal
962. .967
1005. .1010
primer_bind
319 a 279 c 185 g 227 t
BASE COUNT
ORIGIN

alignment_scores:
Quality: 42.00 Length: 8
Ratio: 5.250 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:
US-09-444-281-27 x AMAPID73

Align seg 1/1 to: AMAPID73 from: 1 to: 1010

2 GluA1agluProGluA1agluPro 9
|||||
198 GAAGCTGACGCGAAGCTGAACCC 221

seq_name: gb_in:DDU66367

seq_documentation_block:
LOCUS DDU66367 1390 bp mRNA INV 28-AUG-1996
DEFINITION Dictyostelium discoideum Sapa (sapa) mRNA, partial cds.
ACCESSION U66367
VERSION U66367.1 GI:1513233
KEYWORDS
SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum
Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.
REFERENCE 1 (bases 1 to 1390)
AUTHORS Loomis, W.F.
TITLE Direct Submission
JOURNAL Submitted (08-AUG-1996) Dept. of Biology 0322, University of
California, San Diego, 9500 Gilman Drive, La Jolla, CA 92093-032,
USA

FEATURES
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/chromosome="4"
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gene
CDS

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seq_name: gb_in:DDU72746

seq_documentation_block:
LOCUS DDU72746 1486 bp mRNA INV 19-MAR-1997
DEFINITION Dictyostelium discoideum cysteine proteinase (cprc) mRNA, complete
cds.
ACCESSION U72746
VERSION U72746.1 GI:1644501
KEYWORDS
SOURCE Dictyostelium discoideum.
ORGANISM Dictyostelium discoideum
Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.
REFERENCE 1 (bases 1 to 1486)
AUTHORS Ord, T., Adessi, C., Wang, L. and Freeze, H. H.
TITLE The cysteine proteinase gene cprc in Dictyostelium discoideum has a
serine-rich domain that contains GluNAc-1-P
Arch. Biochem. Biophys. 339 (1), 64-72 (1997)
JOURNAL 97223364
MEDLINE 2 (bases 1 to 1486)
REFERENCE Ord, T., Adessi, C., Wang, L. and Freeze, H. H.
AUTHORS Direct Submission
TITLE Submitted (27-SEP-1996) The Burnham Institute, 10901 N. Torrey
Pines Rd. La Jolla, CA 92093, USA
JOURNAL

FEATURES
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1. .1486
Location/Qualifiers
/organism="Dictyostelium discoideum"
/strain="AX-2"
/db_xref="taxon:44689"
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/protein_id="AAC47482.1"
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AOLSSVNVNNGSGSPDIAAKVTOGPTVAIDASNOFOLYVSGIYNPPASSSTOLDIG
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BASE COUNT 448 a 314 c 278 g 446 t
ORIGIN

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:37 : Search time 24.75 Seconds
(without alignments)
10.001 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAEPKPEPIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 212252 seqs, 22503292 residues

Total number of hits satisfying chosen parameters: 212252

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	39	66.1	486	2 US-08-942-423-3	Sequence 3, Appl1
2	38	64.4	20	1 US-08-208-181A-8	Sequence 8, Appl1
3	38	64.4	20	1 US-08-208-181A-10	Sequence 10, Appl1
4	38	64.4	27	1 US-07-603-782A-1	Sequence 1, Appl1
5	38	64.4	229	2 US-08-394-600B-20	Sequence 20, Appl1
6	38	64.4	229	4 US-08-944-483-30	Sequence 30, Appl1
7	38	64.4	229	5 PCT-US95-02513-20	Sequence 20, Appl1
8	38	64.4	226	2 US-08-230-428B-4	Sequence 4, Appl1
9	37	62.7	226	4 US-08-944-483-41	Sequence 41, Appl1
10	37	62.7	227	4 US-08-944-483-40	Sequence 40, Appl1
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12	37	62.7	247	2 US-08-851-974-1	Sequence 4, Appl1
13	37	62.7	247	2 US-09-213-390-1	Sequence 1, Appl1
14	37	62.7	247	2 US-09-213-390-4	Sequence 4, Appl1
15	37	62.7	248	2 US-08-851-974-3	Sequence 3, Appl1
16	37	62.7	248	2 US-09-213-390-3	Sequence 3, Appl1
17	37	62.7	302	4 US-09-457-046B-18	Sequence 18, Appl1
18	37	62.7	302	3 US-08-996-139-15	Sequence 15, Appl1
19	37	62.7	625	4 US-08-995-659-15	Sequence 15, Appl1
20	37	62.7	625	4 US-09-215-649A-15	Sequence 15, Appl1
21	36	61.0	364	2 US-08-651-940-2	Sequence 2, Appl1
22	36	61.0	364	2 US-09-025-580-37	Sequence 37, Appl1
23	36	61.0	372	4 US-09-286-904-24	Sequence 24, Appl1
24	35	59.3	36	2 US-08-942-423-12	Sequence 12, Appl1
25	35	59.3	433	1 US-08-522-166-7	Sequence 7, Appl1
26	35	59.3	433	1 US-08-488-382A-7	Sequence 7, Appl1
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29	35	59.3	486	2 US-08-942-423-2	Sequence 2, Appl1
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31	34	57.6	10	1 US-08-468-674B-86	Sequence 86, Appl1
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37	34	57.6	59	1 US-08-306-871-25	Sequence 25, Appl1
38	34	57.6	59	1 US-08-569-959-25	Sequence 25, Appl1
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43	34	57.6	326	4 US-09-058-389A-3	Sequence 3, Appl1
44	34	57.6	347	3 US-09-059-369-2	Sequence 2, Appl1
45	34	57.6	456	4 US-09-058-389A-2	Sequence 2, Appl1

ALIGNMENTS

RESULT 1
US-08-942-423-3
; Sequence 3, Application US/08942423
; Patent No. 5891673
GENERAL INFORMATION:
; APPLICANT: Hashimoto, Yasuhiro
; APPLICANT: Takemoto, Yoshihiro
; TITLE OF INVENTION: Lck Binding Protein
; NUMBER OF SEQUENCES: 68
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Syntex (U.S.A.) Inc.
; STREET: 3401 Hillview Ave.
; CITY: Palo Alto
; STATE: California
; COUNTRY: U.S.A.
; ZIP: 94303
COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/942,423
; FILING DATE: 01-Oct-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/362,715
; FILING DATE: 23-DEC-1994
ATTORNEY/AGENT INFORMATION:
; NAME: Perles, Rohan
; REGISTRATION NUMBER: 35,752
; REFERENCE/DOCKET NUMBER: 28260
TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 852-1698
; TELEFAX: (415) 496-3529
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 486 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; HYPOTHEICAL: NO
; ANTI-SENSE: NO
; FRAGMENT TYPE: N-terminal
; ORIGINAL SOURCE:
; ORGANISM: LCK BINDING PROTEIN
; US-08-942-423-3

Query Match 66.1%; Score 39; DB 2; Length 486;
Best Local Similarity 77.8%; Pred. No. 35;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 HEAPEPEEP 9
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Db 360 YEAPPEEP 368

RESULT 2

US-08-208-181A-8
; Sequence 8, Application US/08208181A
; Patent No. 5654167
; GENERAL INFORMATION:
; APPLICANT: Gabay, Joelle E.
; APPLICANT: Nathan, Carl F.
; TITLE OF INVENTION: ANTIMICROBIAL PROTEINS, COMPOSITIONS
; TITLE OF INVENTION: CONTAINING SAME AND USES THEREOF
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 2200 Sand Hill Road, Suite 100
; CITY: Menlo Park
; STATE: CA
; COUNTRY: USA
; ZIP: 94025
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/208,181A
; FILING DATE: 08-MAR-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Bozicevic, Karl
; REGISTRATION NUMBER: 28,807
; REFERENCE/DOCKET NUMBER: 06514/024002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 322-5070
; TELEFAX: (415) 854-0875
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 amino acids
; TYPE: amino acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: protein
; FRAGMENT TYPE: N-terminal
US-08-208-181A-8

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Best Local Similarity 54.5%; Pred. No. 1.6;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

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Db 5 HEAPHSRPM 15

RESULT 3

US-08-208-181A-10
; Sequence 10, Application US/08208181A
; Patent No. 5654167
; GENERAL INFORMATION:
; APPLICANT: Gabay, Joelle E.
; APPLICANT: Nathan, Carl F.
; TITLE OF INVENTION: ANTIMICROBIAL PROTEINS, COMPOSITIONS
; TITLE OF INVENTION: CONTAINING SAME AND USES THEREOF
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 2200 Sand Hill Road, Suite 100
; CITY: Menlo Park
; STATE: CA
; COUNTRY: USA
; ZIP: 94025
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/208,181A
; FILING DATE: 08-MAR-1994
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Bozicevic, Karl
; REGISTRATION NUMBER: 28,807
; REFERENCE/DOCKET NUMBER: 06514/024002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 322-5070
; TELEFAX: (415) 854-0875
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 20 amino acids
; TYPE: amino acid
; STRANDEDNESS: unknown
; TOPOLOGY: unknown
; MOLECULE TYPE: protein
; FRAGMENT TYPE: N-terminal
US-08-208-181A-10

ADDRESSEE: Fish & Richardson P.C.
STREET: 2200 Sand Hill Road, Suite 100
CITY: Menlo Park
STATE: CA
COUNTRY: USA
ZIP: 94025

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/208,181A
FILING DATE: 08-MAR-1994
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Bozicevic, Karl
REGISTRATION NUMBER: 28,807
REFERENCE/DOCKET NUMBER: 06514/024002
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 322-5070
TELEFAX: (415) 854-0875
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 20 amino acids
TYPE: amino acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: protein
FRAGMENT TYPE: N-terminal
US-08-208-181A-10

Query Match 64.4%; Score 38; DB 1; Length 20;
Best Local Similarity 54.5%; Pred. No. 1.6;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

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Db 5 HEAPHSRPM 15

RESULT 4

US-07-603-782A-1
; Sequence 1, Application US/07603782A
; Patent No. 5200319
; GENERAL INFORMATION:
; APPLICANT: Arnaout, Amin M.
; APPLICANT: McCluskey, Robert T.
; APPLICANT: Niles, John L.
; TITLE OF INVENTION: Diagnosis of Glomerulonephritis
; NUMBER OF SEQUENCES: 1
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Fish & Richardson
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb storage
; COMPUTER: IBM PS/2 Model 502 or 55SX
; OPERATING SYSTEM: IBM P.C. DOS (Version 3.30)
; SOFTWARE: WordPerfect (Version 5.0)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/603,782A
; FILING DATE: 19901025
; CLASSIFICATION: 436
; PRIOR APPLICATION DATA:
; PRIOR APPLICATION DATA: including application
; APPLICATION NUMBER: 07/428,286
; FILING DATE: 27-OCT-1989
; ATTORNEY/AGENT INFORMATION:

NAME: Clark, Paul T.
REGISTRATION NUMBER: 30,162
REFERENCE/DOCKET NUMBER: 00786/034002
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 542-5070
TELEFAX: (617) 542-8906
TELEX: 200154
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 27
TYPE: AMINO ACID
TOPOLOGY: linear
US-07-603-782A-1

Query Match 64.4%; Score 38; DB 1; Length 27;
Best Local Similarity 54.5%; Pred. No. 2.3;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
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Db 5 HEAOPHSRPTM 15

RESULT 5
US-08-394-600B-20
Sequence 20, Application US/08394600B
Patent No. 5843693
GENERAL INFORMATION:
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Kochs, Kirsten E.
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses Thereof
NUMBER OF SEQUENCES: 28
CORRESPONDENCE ADDRESS:
ADDRESSEE: McAndrews, Held & Malloy, Ltd.
STREET: 500 West Madison Street, 34th Floor
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60661
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: EC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/394,600B
FILING DATE: 02/27/95
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Donald J. Pochopien
REGISTRATION NUMBER: 32,167
REFERENCE/DOCKET NUMBER: 820,005/11850US05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/707-8889
TELEFAX: 312/707-9155
TELEX:
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-394-600B-20

Query Match 64.4%; Score 38; DB 2; Length 229;
Best Local Similarity 54.5%; Pred. No. 2.3;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
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Db 5 HEAOPHSRPTM 15

RESULT 6
US-08-944-483-30
Sequence 30, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLPITTS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLASS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: OF THE PROSTATE
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM compatible
OPERATING SYSTEM: DOS
SOFTWARE: FASTSEQ for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ ID NO: 30:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: NO. 6232456e
US-08-944-483-30

Query Match 64.4%; Score 38; DB 4; Length 229;
Best Local Similarity 54.5%; Pred. No. 2.3;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
|||:|:|
Db 5 HEAOPHSRPTM 15

RESULT 7
PCT-US95-02513-20
Sequence 20, Application PC/TUS9502513
GENERAL INFORMATION:

APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Koths, Kirston E.
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
TITLE OF INVENTION: Compositions for the inhibition of
TITLE OF INVENTION: Protein Hormone Formation and Uses thereof
NUMBER OF SEQUENCES: 22
CORRESPONDENCE ADDRESS:
ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
ADDRESSEE: Borun
STREET: 6300 Sears Tower, 233 South Wacker Drive
CITY: Chicago
STATE: Illinois
COUNTRY: United States of America
ZIP: 60606-6402
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/02513
FILING DATE:
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Williams Jr., Joseph A.
REGISTRATION NUMBER: 38,659
REFERENCE/DOCKET NUMBER: 27527/32404
TELECOMMUNICATION INFORMATION:
TELEPHONE: 312/474-6960
TELEFAX: 312/474-0448
TELEX: 25-3856
INFORMATION FOR SEQ ID NO: 20:
SEQUENCE CHARACTERISTICS:
LENGTH: 229 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
PCT-US95-02513-20

Query Match 64.4% Score 38; DB 5; Length 229;
Best Local Similarity 54.5% Pred. No. 23;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
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DB 5 HEAPHHSRPYM 15

RESULT 8
US-08-230-428B-4
Sequence 4, Application US/08230428B
Patent No. 5998378
GENERAL INFORMATION:
APPLICANT: Kriegluer, Michael
APPLICANT: Perez, Carl
APPLICANT: Halenbeck, Robert F.
APPLICANT: Jewell, David A.
APPLICANT: Koths, Kirston E.
TITLE OF INVENTION: Compositions For the Inhibition Of TNF
TITLE OF INVENTION: Hormone Formation And Uses Thereof (As Amended)
NUMBER OF SEQUENCES: 14
CORRESPONDENCE ADDRESS:
ADDRESSEE: CHIRON CORPORATION Intellectual Property - R440
STREET: 4560 Horton Street, P.O. Box 8097
CITY: Emeryville
STATE: California
COUNTRY: United States of America
ZIP: 94662-8097
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/230,428B
FILING DATE: 19-APR-1994
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/905,546
FILING DATE: 25-JUN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 07/395,253
FILING DATE: 16-AUG-1989
ATTORNEY/AGENT INFORMATION:
NAME: Saveriede, Paul B.
REGISTRATION NUMBER: 36,914
REFERENCE/DOCKET NUMBER: 0820,004
TELECOMMUNICATION INFORMATION:
TELEPHONE: (510) 601-2718
TELEFAX: (510) 655-3542
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 256 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-230-428B-4

Query Match 64.4% Score 38; DB 2; Length 256;
Best Local Similarity 54.5% Pred. No. 26;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPIM 11
|||:|:|
DB 32 HEAPHHSRPYM 42

RESULT 9
US-08-944-483-41
Sequence 41, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLPITTS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLASS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
APPLICANT: STROUPE, STEVEN D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: AND METHODS USEFUL FOR DETECTING AND TREATING DISEASES
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:

FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ. ID NO: 41:
SEQUENCE CHARACTERISTICS:
LENGTH: 226 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: No. 6232456e
US-08-944-483-41

Query Match 62.7%; Score 37; DB 4; Length 226;
Best Local Similarity 54.5%; Pred. No. 33;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPM 11
|||:|:|
DB 5 HEAKPHSRPYM 15

RESULT 10
US-08-944-483-40
Sequence 40, Application US/08944483
Patent No. 6232456
GENERAL INFORMATION:
APPLICANT: COHEN, MAURICE
APPLICANT: COLPITS, TRACEY L.
APPLICANT: FRIEDMAN, PAULA N.
APPLICANT: GRANADOS, EDWARD N.
APPLICANT: KLAS, MICHAEL R.
APPLICANT: RUSSELL, JOHN C.
APPLICANT: STEWART, KENT D.
APPLICANT: STROUPE, STEVEN D.
TITLE OF INVENTION: NOVEL SERINE PROTEASE REAGENTS
TITLE OF INVENTION: AND METHODS USEFUL FOR DETECTING AND TREATING DISEASES
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Abbott Laboratories
STREET: 100 Abbott Park Road
CITY: Abbott Park
STATE: IL
COUNTRY: USA
ZIP: 60064-3500
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/944,483
FILING DATE:
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Becker, Cheryl L.
REGISTRATION NUMBER: 35,441
REFERENCE/DOCKET NUMBER: 6183.US.01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 847/935-1729
TELEFAX: 847/938-2623
TELEX:
INFORMATION FOR SEQ. ID NO: 40:

SEQUENCE CHARACTERISTICS:
LENGTH: 227 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: No. 6232456e
US-08-944-483-40

Query Match 62.7%; Score 37; DB 4; Length 227;
Best Local Similarity 54.5%; Pred. No. 33;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPM 11
|||:|:|
DB 5 HEAKPHSRPYM 15

RESULT 11
US-08-851-974-1
Sequence 1, Application US/08851974
Patent No. 5858758
GENERAL INFORMATION:
APPLICANT: Hillman, Jennifer L.
APPLICANT: Corley, Neil C.
APPLICANT: Shah, Puryi
TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
NUMBER OF SEQUENCES: 4
CORRESPONDENCE ADDRESS:
ADDRESSEE: Incyte Pharmaceuticals, Inc.
STREET: 3174 Porter Drive
CITY: Palo Alto
STATE: CA
COUNTRY: USA
ZIP: 94304
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq for Windows Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/851,974
FILING DATE: Filed Herewith
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0288 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
INFORMATION FOR SEQ. ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 247 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: NCBI
CLONE: 854243
US-08-851-974-1

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

RESULT 12
US-08-851-974-4
; Sequence 4, Application US/08851974
; Patent No. 5858758
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; FILING DATE: Filed Herewith
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 306682
; US-08-851-974-4

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

RESULT 13
US-09-213-390-1
; Sequence 1, Application US/09213390
; Patent No. 5965711
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA

ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/213,390
; FILING DATE:
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/851,974
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 247 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: NCANOT01
; CLONE: 854243
; US-09-213-390-1

Query Match 62.7%; Score 37; DB 2; Length 247;
Best Local Similarity 54.5%; Pred. No. 36;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
|||:|:|
DB 25 HEAKPHSRPYM 35

RESULT 14
US-09-213-390-4
; Sequence 4, Application US/09213390
; Patent No. 5965711
; GENERAL INFORMATION:
; APPLICANT: Hillman, Jennifer L.
; APPLICANT: Corley, Neil C.
; APPLICANT: Shah, Purvi
; TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Incyte Pharmaceuticals, Inc.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/213,390
; FILING DATE:
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/851,974
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0288 US
; TELECOMMUNICATION INFORMATION:

TELEPHONE: 415-855-0555
 TELEFAX: 415-845-4166
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 247 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 IMMEDIATE SOURCE:
 LIBRARY: GenBank
 CLONE: 306682
 US-09-213-390-4

QY 1 HEAPEAPEPTM 11
 |||:|:|
 Db 25 HEAKPHSRPYM 35
 Search completed: January 4, 2002, 08:40:57
 Job time: 140 sec

Query Match 62.7%; Score 37; DB 2; Length 247;
 Best Local Similarity 54.5%; Pred. No. 36;
 Matches 6; Conservat. 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAPEPTM 11
 |||:|:|
 Db 25 HEAKPHSRPYM 35

RESULT 15
 US-08-851-974-3
 Sequence 3, Application US/08851974
 Patent No. 5858758
 GENERAL INFORMATION:
 APPLICANT: Hillman, Jennifer L.
 APPLICANT: Corley, Neil C.
 TITLE OF INVENTION: HUMAN SERINE PROTEASE PRECURSOR
 NUMBER OF SEQUENCES: 4
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Incyte Pharmaceuticals, Inc.
 STREET: 3174 Porter Drive
 CITY: Palo Alto
 STATE: CA
 COUNTRY: USA
 ZIP: 94304
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: DOS
 SOFTWARE: FASTSEQ for Windows Version 2.0
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/851,974
 FILING DATE: Filed Herewith
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER:
 FILING DATE:
 ATTORNEY/AGENT INFORMATION:
 NAME: Billings, Lucy J.
 REGISTRATION NUMBER: 36,749
 REFERENCE/DOCKET NUMBER: PF-0288 US
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 415-855-0555
 TELEFAX: 415-845-4166
 INFORMATION FOR SEQ ID NO: 3:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 248 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 IMMEDIATE SOURCE:
 LIBRARY: GenBank
 CLONE: 206690
 US-08-851-974-3

Query Match 62.7%; Score 37; DB 2; Length 248;
 Best Local Similarity 54.5%; Pred. No. 36;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:38:37 ; Search time 53.46 Seconds
(without alignments)
15.241 Million cell updates/sec

Title: US-09-444-281-27
Perfect score: 59
Sequence: 1 HEAEPFAEPIM 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 522463 seqs, 74073290 residues
Total number of hits satisfying chosen parameters: 522463

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

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2: /SID58/gcgdata/geneseq/geneseq/AA1981.DAT:*

3: /SID58/gcgdata/geneseq/geneseq/AA1982.DAT:*

4: /SID58/gcgdata/geneseq/geneseq/AA1983.DAT:*

5: /SID58/gcgdata/geneseq/geneseq/AA1984.DAT:*

6: /SID58/gcgdata/geneseq/geneseq/AA1985.DAT:*

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21: /SID58/gcgdata/geneseq/geneseq/AA2000.DAT:*

22: /SID58/gcgdata/geneseq/geneseq/AA2001.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	39	66.1	6797	22	AA831558 Pimaricin biosynth
2	38	64.4	27	12	AA811996 N-terminal of p29
3	38	64.4	27	14	AA834444 N-terminus of p29
4	38	64.4	237	13	AA820509 Human proteinase 3
5	38	64.4	236	16	AA845403 Deduced sequence o
6	38	64.4	256	16	AA85639 MY17 preproPR-3.
7	37	62.7	149	22	AAU00451 Protein encoded by
8	37	62.7	173	22	AA872305 Sunflower glutamic
9	37	62.7	246	12	AA813253 Human cytotoxic ce
10	37	62.7	247	20	AA84158 Human serine pro
11	37	62.7	261	19	AA854091 Homo sapiens BR2 s

12	37	62.7	267	21	AA90291 Human peptidase, H
13	37	62.7	267	22	AA820156 Human protein SECP
14	37	62.7	278	21	AA858142 Lung cancer associ
15	37	62.7	302	22	AAE00235 Protein encoded by
16	37	62.7	553	19	AA863703 Truncated rat RSK3
17	37	62.7	567	20	AA868788 Polyketide fragme
18	37	62.7	568	21	AA85658 Human Acinus S pro
19	37	62.7	571	20	AA841704 Human PRO351 prote
20	37	62.7	571	21	AA84260 Human PRO351 (UNO3
21	37	62.7	571	21	AA824046 Human PRO351 prote
22	37	62.7	583	21	AA85659 Human Acinus S' pr
23	37	62.7	625	19	AA863200 Murine osteoclast
24	37	62.7	625	19	AA869558 Murine NF-KB recep
25	37	62.7	625	19	AA868294 Murine NF-KB recep
26	37	62.7	625	21	AA859509 OBM binding protei
27	37	62.7	625	21	AA83649 A mouse receptor a
28	37	62.7	625	22	AA804427 Murine receptor ac
29	37	62.7	625	22	AAE01994 Murine RANK (recep
30	37	62.7	732	19	AA863715 Rat RSK3 protein.
31	37	62.7	1341	21	AA85657 Human Acinus L pro
32	37	62.7	75	21	AA802199 Human secreted pro
33	36	61.0	108	21	AA833906 Human secreted pro
34	36	61.0	161	21	AA842933 Arabidopsis thalia
35	36	61.0	181	21	AA817997 Arabidopsis thalia
36	36	61.0	181	21	AA847650 Arabidopsis thalia
37	36	61.0	183	21	AA841769 Human OREF ORF153
38	36	61.0	192	22	AA814154 Human novel protei
39	36	61.0	196	21	AA842932 Arabidopsis thalia
40	36	61.0	213	21	AA843275 Human OREF ORF3039
41	36	61.0	216	21	AA817996 Arabidopsis thalia
42	36	61.0	216	21	AA847649 Arabidopsis thalia
43	36	61.0	252	21	AA815889 Arabidopsis thalia
44	36	61.0	254	21	AA847624 Arabidopsis thalia
45	36	61.0	264	21	AA847623 Arabidopsis thalia

ALIGNMENTS

RESULT 1	
AA831558	standard; Protein: 6797 AA.
ID	AA831558
AC	AA831558
XX	
XX	20-APR-2001 (first entry)
DE	Pimaricin biosynthesis associated polyketide synthase polypeptide.
DE	Polyketide synthase; oxidative modification; metabolite; antibiotic;
KW	antibiotic; pimaricin.
KW	antibiotic; pimaricin.
XX	
OS	Streptomyces natalensis.
XX	
XX	WO200077222-A1.
PN	
XX	21-DEC-2000.
PD	
XX	14-JUN-2000; 2000WO-EP06227.
PF	
XX	14-JUN-1999; 99EP-0201893.
PR	
XX	(STAM) DSM NV.
PA	
XX	Martin JF, Aparicio JF, Collina AJ;
PI	
XX	WPI; 2001-080693/09.
XX	DR
XX	New polynucleotides encoding enzymes involved in the biosynthesis of
XX	pimaricin, useful for modifying the biosynthesis of pimaricin and in
PT	the synthesis of new compounds
PT	
XX	

PS Disclosure; Page 81-101; 116pp; English.

XX The present sequence represents a polyketide synthase which is associated
 CC with the biosynthesis of pimarinin. The polyketide synthase polyketide
 CC is useful for the oxidative modification of a methyl group of a suitable
 CC compound, e.g. a bioactive compound including a secondary metabolite,
 CC antibiotics and anticancer agents. Recombinant cells comprising the
 CC gene are useful for the production of pimarinin. The polyketide synthase
 CC polynucleotide may be over expressed in Streptomyces, leading to an
 CC increase in the biosynthesis of pimarinin, as a source of primers for
 CC amplification reaction and as probes.

XX Sequence 6797 AA:

XX SQ

Query Match 66.1%; Score 39; DB 22; Length 6797;
 Best Local Similarity 77.6%; Pred. No. 1.5e+03;
 Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 HEAPEAP 9
 |||||
 Db 463 heapeap 471

RESULT 2
 AAR1996
 ID AAR1996 standard; peptide; 27 AA.
 XX
 AC AAR1996;
 XX
 DT 26-JUL-1991 (first entry)
 XX
 DE N-terminal of p29 protein.
 XX
 XX Wegener's granulomatosis; monoclonal antibodies; autoantibodies;
 KM glomerulonephritis; serine protease; antigen.
 XX
 OS Homo sapiens.
 XX
 PN W09106572-A.
 XX
 PD 16-MAY-1991.
 XX
 PF 29-OCT-1990; 90MO-US06277.
 XX
 PR 27-OCT-1989; 89US-0428286.
 XX
 PA (GEHO-) GEN HOSPITAL CORP.
 XX
 PI Arnaout M, McCluskey RT, Niles J;
 XX
 DR WPI; 1991-164137/22.
 XX
 XX Purified p29 protein - used to detect auto-antibodies diagnostic
 PT for Wegener's granulomatosis and conditions associated with
 PT glomerulo nephritis
 XX
 XX Claim 1; Page 22; 33pp; English.

XX The p29 protein is a 29 kD antigen which was prepd. by affinity
 CC purification from neutrophil acid extract, using 1B8 monoclonal
 CC antibodies. The purified antigen migrated on SDS-PAGE as three
 CC close bands, with major component at 29 kD under non-reducing
 CC conditions. It reacted with autoantibodies from patients sera
 CC indicating identity between Wegener's granulomatosis autoantigen.
 CC On isofocusing gels it had a pI of 9.2-9.4. It is a novel serine
 CC protease showing homology with leukocyte elastase and cathepsin G.
 CC The protein or monoclonal Abs can be used to test for the presence
 CC of autoantibodies diagnostic for Wegener's granulomatosis. In
 CC combination with myeloperoxidase (and/or Abs against it), p29 Mabs
 CC can also be used to test for autoantibodies associated with pauci
 CC immune necrotizing and/or crescentic glomerulonephritis.

SQ Sequence 27 AA:

XX Query Match 64.4%; Score 38; DB 12; Length 27;
 Best Local Similarity 54.5%; Pred. No. 7.7;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPIM 11
 |||||
 Db 5 heapshsrypm 15

RESULT 3
 AAR34444
 ID AAR34444 standard; peptide; 27 AA.
 XX
 AC AAR34444;
 XX
 DT 29-JUL-1993 (first entry)
 XX
 DE N-terminus of p29.
 XX
 XX Pauci-immune necrotizing; crescentic glomerulonephritis; neutrophil;
 KM autoantibodies; Wegener's granulomatosis; microscopic polyarteritis
 KM nodosa; primary MCGN.
 XX
 OS Homo sapiens.
 XX
 PN US5200319-A.
 XX
 PD 06-APR-1993.
 XX
 PF 27-OCT-1989; 89US-0428286.
 XX
 PR 27-OCT-1989; 89US-0428286.
 XX
 PR 25-OCT-1990; 90US-0603782.
 XX
 PA (GEHO-) GEN HOSPITAL CORP.
 XX
 PI Arnaout MA, McCluskey RT, Niles JL;
 XX
 DR WPI; 1993-133732/16.
 XX
 XX Diagnosis of pauci-immune necrotizing and/or crescentic
 PT glomerulonephritis - using p29 protein isolated from
 PT neutrophil(s), and which binds auto-antibodies present in serum
 PT of patients with Wegener's granulomatosis
 XX
 PS Disclosure; Page 12; 13pp; English.

XX The p29 protein can be isolated from neutrophils, has a mol. wt. of
 CC 29 kD (SDS PAGE), binds disopropyl fluorophosphate, has a pI of 9.2-
 CC 9.4, is capable of binding auto antibodies present in the sera of
 CC individuals afflicted with Wegener's granulomatosis and has the N-
 CC terminal sequence shown. The p29 protein can be used for diagnosis
 CC of Wegener's granulomatosis, microscopic polyarteritis nodosa and
 CC primary MCGN.
 XX
 XX Sequence 27 AA:

XX SQ

Query Match 64.4%; Score 38; DB 14; Length 27;
 Best Local Similarity 54.5%; Pred. No. 7.7;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEAPIM 11
 |||||
 Db 5 heapshsrypm 15

RESULT 4
 AAR20509
 ID AAR20509 standard; protein; 237 AA.

```

XX AC AAR20509;
XX XX
XX DT 06-MAY-1992 (first entry)
XX DE Human proteinase 3.
XX KM Serine protease; polymerase chain reaction; PCR; autoantibody.
XX OS Homo sapiens.
XX FH Key
XX FT Peptide
XX FT 1..6
XX FT /label= signal
XX FT /note= "part of signal peptide"
XX FT Peptide
XX FT 7..8
XX FT /label= propeptide
XX FT Protein
XX FT 9..237
XX FT /label= proteinase_3
XX PN MO9200378-A.
XX PD 09-JAN-1992.
XX PF 20-JUN-1991; 91MO-1001142.
XX PR 22-JUN-1990; 90DE-4019984.
XX PA (GBFB-) GBF GES BIOTECHN EQ.
XX PI Jenne D, Tschopp J, Ludemann J, Utecht B, Gross LB;
XX DR WPI: 1992-041560/05.
XX DR N-PSDB; AAQ20727.
XX PT DNA sequence coding for proteinase 3 - allows mass prodn. of
XX PT enzyme, with modifications for improved therapy and diagnosis of
XX PT Wegener's granulomatosis
XX PS Disclosure: Fig 3; 21pp; German.
XX CC The CDNA sequence corresponding to the coding region of the
XX CC proteinase 3 gene was obtained by PCR amplification of cDNA from
XX CC cell line U937 (see AAQ22231 and AAQ22232 for PCR primer sequences). An
XX CC amplified product of 784bp was obtained. The product was cut with
XX CC EcoRI and cloned into EcoRI-digested M13mp18/BAP. The complete
XX CC sequence was determined, and the amino acid sequence it encodes was
XX CC deduced from it.
XX SQ Sequence 237 AA;

Query Match 64.4%; Score 38; DB 13; Length 237;
Best Local Similarity 54.5%; Pred. No. 71;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
DB 13 heagphsrypm 23

RESULT 5
AAR45403
ID AAR45403 standard; Protein; 256 AA.
AC AAR45403;
XX AC
XX DT 18-JUL-1994 (first entry)
XX DE Deduced sequence of human proteinase-3 (PR-3).
XX KM TNF convertase; proteinase-3; PR-3; tumour necrosis factor;
XX KM protNF.

```

```

XX OS Homo sapiens.
XX XX
XX PN MO9400555-A.
XX XX
XX PD 06-JAN-1994.
XX PF 25-JUN-1993; 93MO-US06120.
XX PR 25-JUN-1992; 92US-0905546.
XX PA (CETU ) CETUS ONCOLOGY CORP.
XX FH Halenbeck RF, Jewell DA, Kothe KE, Kriegler M, Perez C;
XX FT WPI: 1994-026195/03.
XX FT N-PSDB; AAQ54498.
XX DR
XX PT Cgds. which inhibit formation of mature tumour necrosis factor
XX PT from its precursor - identified using TNF convertase, e.g.
XX PT mutein(s), antibodies or peptide phosphonate(s), for preventing
XX PT and treating sepsis, AIDS, auto-immune disease etc.
XX PS Disclosure: Fig 2; 69pp; English.
XX CC PROTNF refers to TNF having a molecular weight of about 26,000,
XX CC which is the prohormone form of TNFa. PROTNF is cleaved to a lower
XX CC molecular weight 'mature' form, pref. 17kD, which, in its multimeric
XX CC (usually trimeric) form, is substantially involved in producing life-
XX CC threatening physiological changes associated with sepsis. PROTNF is
XX CC cleaved by convertase. One TNF convertase is serine protease
XX CC proteinase-3, also called PR-3, P-29B or myeloblastin. A suitable
XX CC source of convertase is the HL60 cell line (or extracts, or the
XX CC culture media in which it is grown). The convertase produced by
XX CC HL60 has been sequenced and is identical to the known lymphocyte
XX CC serine protease PR-3 which has other activities unrelated to TNF
XX CC processing.
XX SQ Sequence 256 AA;

Query Match 64.4%; Score 38; DB 15; Length 256;
Best Local Similarity 54.5%; Pred. No. 76;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 HEAPEAEPIIM 11
DB 32 heagphsrypm 42

RESULT 6
AAR85639
ID AAR85639 standard; Protein; 256 AA.
AC AAR85639;
XX AC
XX DT 23-APR-1996 (first entry)
XX DE MY17 preproPR-3.
XX KM PR-3; preproPR-3; MY17; human neutrophil protease-3; serine protease;
XX KM tumour necrosis factor alpha; TNFalpha; HL60; MY17; B cell; T cell;
XX KM tumour necrosis factor alpha convertase; cytokine; septic shock;
XX KM rheumatoid arthritis; cachexia; cerebral malaria; graft-host disease;
XX KM ischaemia/reperfusion injury; autoimmune disease; AIDS.
XX OS Homo sapiens.
XX FH Key
XX FT Peptide
XX FT 1..25
XX FT /note= "leader sequence present only in preproPR-3"
XX FT Peptide
XX FT 26..27
XX FT /note= "dipeptide present in proPR-3"

```

FT	Protein	28.256	/note="mature PR-3"
XX			
PN	WO9524501-A1.		
XX			
PD	14-SEP-1995.		
XX			
PF	02-MAR-1995;	95MO-US02513.	
PR	28-FEB-1995;	95US-0395456.	
PR	07-MAR-1994;	94US-0208574.	
PR	19-APR-1994;	94US-0230428.	
XX	27-FEB-1995;	95US-0394600.	
PA	(CETU) CETUS ONCOLOGY CORP.		
PI	Halenback RF, Jewell DA, Kolts KE, Krieglger M, Perez C;		
DR	WPI; 1995-328287/42.		
N-PSDB; AAT02565.			
PT	Identification of inhibitors of mature TNFalpha prodn. - useful for treatment of septic shock, rheumatoid arthritis, etc..		
XX			
PS	Example 2; Page 82; 96pp; English.		
CC	This sequence represents the preproPR-3. PR-3 is active recombinant human neutrophil protease-3. PR-3 is a serine protease, and is a tumour necrosis factor alpha (TNFalpha) convertase. The cDNA encoding this sequence was isolated from the H60 cell clone MY17. The mature PR-3 can be used in the method of the invention for identifying agents that inhibit cleavage of proTNFalpha to mature TNFalpha. In the method, proTNFalpha is incubated with PR-3 (or another TNFalpha convertase), and the cleavage of the proTNFalpha is measured by a colourimetric assay. This is then repeated in the presence of a test compound that is thought to inhibit this process. The results of the two reactions are then compared to determine whether the test compound is an inhibitor. The cleavage inhibitors can be used in the treatment of septic shock, rheumatoid arthritis, cachexia, cerebral malaria, ischaemia/reperfusion injury, graft-host disease, autoimmune diseases, and AIDS. PR-3 can be used to treat unwanted B cell/T cell interaction by contacting it with T cells to cause the release of membrane-bound cytokines.		
XX			
SQ	Sequence	256 AA;	
	Query Match	64.4%;	Score 38; DB 16; Length 256;
	Best Local Similarity	54.5%;	Pred. No. 76;
	Matches	6; Conservative	2; Mismatches
OY	1 HEAEPAEIPIM 11		Indels
	:		Gaps
Db	32 headphsrpyrn 42		
RESULT	7		
AU000451	ID	AU000451 standard; Protein; 149 AA.	
XX			
AC	AU000451;		
XX			
DT	29-MAY-2001	(first entry)	
XX			
DE	Protein encoded by sugarcane promoter cDNA clones c51, c511 and c512.		
KM	Sugarcane promoter region; monocotyledonous plant; stem tissue;		
KM	insecticide; herbicide; disease resistance; improved food content;		
KM	beta-glucuronidase; GUS; starch biosynthesis; fatty acid biosynthesis;		
KM	ADP-glucose pyrophosphorylase; sucrose metabolism; clone c51; c511; c512.		
XX			
OS	Saccharum sp.		
XX			
RH-	Key	Location/Qualifiers	

```

FT      Misc-difference 41
FT      /note= "Encoded by ACG in cDNA clone c51 (AAS01022)"
FT
FT      Misc-difference 92
FT      /label= Ala, Val
FT      /note= "Residue 92 is Ala as deduced from cDNA clones
FT      c51 and c511 (AAS01022-AAS01023) and Val as
FT      deduced from clone c512 (AAS01024)"
FT
FT      Misc-difference 103
FT      /note= "Encoded by TAG in cDNA clone c511 (AAS01023)"
FT
FT      Misc-difference 125
FT      /label= Val, Ala
FT      /note= "Residue 125 is Val as deduced from cDNA clone
FT      c51 (AAS01022) and Ala as deduced from clones
FT      c511 and c512 (AAS01023-AAS01024)"
FT
FT      Misc-difference 130
FT      /note= "Encoded by TCC in cDNA clone c511 (AAS01023)"
FT
FT      Misc-difference 135
FT      /note= "Encoded by GCC in cDNA clone c511 (AAS01023)"
FT
XX      WO200118211-A1.
XX
XX      15-MAR-2001.
XX
XX      PD
XX      01-SEP-2000; 2000WO-AU01033.
XX
XX      PR      02-SEP-1999; 99AU-0002625.
XX
XX      PA      (UYOU) UNIV QUEENSLAND.
XX
XX      PI      Potler B, Birch RG;
XX      PT      WPI: 2001-218560/22.
XX      DR      N-PSDB; AAS01022, AAS01023, AAS01024.
XX
XX      New sugarcane plant promoters for directing expression of heterologous
XX      nucleic acids in a constitutive or tissue-specific manner in
XX      monocotyledonous plants
XX
XX      Claim 40; Fig 15; 107pp; English.
XX
XX      The present sequence represents the polypeptide encoded by sugarcane
XX      plant promoter cDNA isolated from clones c51, c511 and c512. Clone c51,
XX      c511 and c512 promoter cDNAs are 3 of 11 promoter regions of a
XX      transcribable DNA sequence isolated from various sugarcane cDNA clones
XX      (AAS01021-AAS01031). Also described are 4 promoter regions of specific
XX      transcribed DNA sequences (AAS01032-AAS01035). The nucleic acids are
XX      useful for producing transgenic plants, having an altered phenotype and
XX      for driving expression of a foreign or endogenous DNA sequence, which
XX      encode agronomic properties including insecticide, herbicide, disease
XX      resistance, stress tolerance and improved food content, or increased
XX      yields. The foreign or endogenous DNA sequence may comprise a region
XX      transcribed into an antisense RNA or ribozyme that modulates the
XX      expression of a corresponding target gene, or it may encode
XX      beta-glucuronidase (GUS), luciferase, neomycin phosphotransferase, a
XX      product conferring herbicide tolerance, a product affecting starch
XX      biosynthesis or modification, ADP-glucose pyrophosphorylase, a product
XX      involved in fatty acid biosynthesis, a product conferring insect
XX      resistance, a product altering sucrose metabolism or a gene encoding
XX      valuable pharmaceuticals, e.g. antibiotics, secondary metabolites or
XX      vaccines. The promoters are capable of directing high level expression in
XX      many or all cells of a plant, preferentially in stem or meristem tissue
XX      of monocotyledonous plants.
XX
XX      Sequence 149 AA;
XX
XX      Query Match 62.7%; Score 37; DB 22; Length 149;
XX      Best Local Similarity 100.0%; Pred. No. 64;
XX      Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX      3 APEAP 9
XX      |||||
XX      69 aapeap 75

```

DB	76	gapopen	83
OY	2	EAAPEAAP	9
	1		
Query Match	62.7%	Score 37:	DB 22: Length 173;
Best Local Similarity	87.5%	Pred. No. 75:	
Matches	7:	Conservative	0: Mismatches 1: Indels 0 Gaps 0

RESULT	9	
ID	AA13253	standard; Protein: 246 AA.
AC	AA13253;	
DT	11-OCT-1991	(first entry)
DE	Human Cytotoxic Cell Protease-X.	
KW	hccpx inhibitor; cytotoxic T-lymphocytes; CTL.	
OS	Homo sapiens.	
PN	W09110685-A.	
PD	25-JUL-1991.	
PF	17-JAN-1991;	91WO-US00340.
PR	19-JAN-1990;	90US-0467880.
PA	(SERA-) SERAGEN INC.	
PI	Blackley RC, Lobe CG, Paetkau VH, James MN, Murphy M;	
DR	WPI; 1991-237989/32.	
DR	N-PSDB; AAQ12865, AAQ12866.	
PT	DNA vectors, and inhibitors of cytotoxic cell protease - for	
PT	treatment of auto-immune diseases e.g. pernicious anaemia,	
PT	rheumatoid arthritis, allo-graft rejection etc.	
PS	Claim 8; Fig 9; 62pp; English.	
CC	The hccpx gene was isolated from cytotoxic T-cell lymphocytes using	
CC	murine CCP cDNA sequences as probes. The cDNA sequence (i.e. exons	
CC	only) was deduced from the genomic sequence. This is the amino acid	
CC	sequence deduced from the coding regions. A structural analysis of	
CC	the deduced protein was used to design competitive inhibitors of the	
CC	protease. See also AAQ12862-5 and AA13254-R13262.	
SO	Sequence	246 AA;
QY	1 HEAPEAEPIIM 11	
DB	25 heakphsrypym 35	
RESULT	10	
ID	AAW84158	
AC	AAW84158	standard; Protein: 247 AA.
DT	25-FEB-1999	(first entry)
DE	A human serine protease precursor (HSP) protein.	
KW	Human serine protease precursor; HSP; incyte clone 854243;	
KW	immunological disorder; cancer; antagonist; allergy; asthma;	
KW	Crohn's disease; multiple sclerosis; rheumatoid arthritis; glioma;	
OS	lymphoma; myeloma.	
OS	Homo sapiens.	
FT	key	Location/Qualifiers
FT	Modified-site	71

FT Modified-site /note= "potential N-linked glycosylation site"
 FT 104 /note= "potential N-linked glycosylation site"
 FT Disulfide-bond 49 /note= "potential disulphide bridging site"
 FT Disulfide-bond 65 /note= "potential disulphide bridging site"
 FT Disulfide-bond 142 /note= "potential disulphide bridging site"
 FT Disulfide-bond 173 /note= "potential disulphide bridging site"
 FT Disulfide-bond 188 /note= "potential disulphide bridging site"
 FT Disulfide-bond 209 /note= "potential disulphide bridging site"
 FT Disulfide-bond 230 /note= "potential disulphide bridging site"
 FT Disulfide-bond 230 /note= "potential disulphide bridging site"
 FT Region 37..48 /note= "potential substrate binding region"

PN W09850424-A2.

PD 12-NOV-1998.

PF 06-MAY-1998; 98MO-US09096.

PR 07-MAY-1997; 97US-0851974.

PA (INCY-) INCYTE PHARM INC.

PI Corley NC, Hillman JL, Shah P;

DR WPI; 1999-034707/03.

DR N-PSDB; AAV82707.

XX New human serine protease precursor and related nucleic acid.
 PT vectors - for treatment, prevention and diagnosis of immunological
 PT disorders and cancer

PS Claim 1: Fig 1A-C; 56pp; English.

CC The present sequence represents a human serine protease precursor (HSP)
 CC protein. Nucleic acids encoding HSP were first identified in incyte
 CC clone 854243 from the ganglioneuroblastoma tissue CDNA library. The protein
 CC has homology with the rat natural killer cell protease-1 precursor
 CC (RNPR-1) and a human serine protease from cytotoxic T lymphocytes (SECT).
 CC As increased levels of the protein are associated with immunological
 CC disorders and cancers, antagonists are used to treat or prevent
 CC conditions such as allergy, asthma, Crohn's disease, multiple sclerosis,
 CC rheumatoid arthritis, glioma, lymphoma, myeloma etc. The HSP nucleic
 CC acid sequence and its fragments are used as antisense/ribozyme
 CC therapeutics, for detecting and quantifying gene expression (as probes
 CC and primers in standard hybridisation and amplification assays), and for
 CC isolating related sequences and for chromosome mapping.

XX Sequence 247 AA;

Query Match 62.7%; Score 37; DB 20; Length 247;
 Best Local Similarity 54.5%; Pred. No. 1.1e+02;

Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 HEAPEPEPIM 11
 DB 25 heakphsrypm 35

RESULT 11

ID AAM54091 standard; Protein; 261 AA.

XX AC AAM54091;
 XX XX

DT 28-SEP-1998 (first entry)

XX Homo sapiens BE2 sequence.

DE BARD1; ring protein; BRCA1; breast cancer; risk; diagnosis.

XX Homo sapiens.

OS Key Location/Qualifiers

FT Misc-difference 104 /label= Ala, Ser, Pro, Thr

FT Misc-difference 114 /label= Gly

FT Misc-difference 218 /label= Ala

XX W09812327-A2.

PD 26-MAR-1998.

PF 19-SEP-1997; 97WO-US16842.

PR 04-APR-1997; 97US-0042985.

PR 20-SEP-1996; 96US-0025296.

PR 03-APR-1997; 97US-0042611.

PA (TEXA) UNIV TEXAS SYSTEM.

PI Baer R, Bowcock AM;

DR WPI; 1998-230317/20.

DR N-PSDB; AAV24126.

XX DNA sequence encoding BARD1, B123, BE2, BE14, BE31 or BE445 - which
 PT as breast cancer antigen, BRCA1, binding proteins are useful to
 PT identify patient having or at risk of developing cancer

XX Disclosure: Page 272-273; 348pp; English.

CC The sequence is that of a protein which can be used in the
 CC preparation of the recombinant breast cancer antigen, BRCA1, binding
 CC proteins BARD1, B123, BE2, BE14, BE31 or BE445, or a composition for the
 CC detection of a BARD1, B123, BE2, BE14, BE31 or BE445 nucleic acid
 CC sequence, specifically a wild type BARD1 composition for the detection
 CC or purification of BRCA1, useful to identify a patient having, or at
 CC risk of developing cancer. BARD1 can be used in the preparation of an
 CC anti-BARD1 antibody, and in the detection and purification of a BRCA1
 CC protein. BARD1, B123, BE2, BE14, BE31 or BE445 can be used in the
 CC identification of a binding protein agonist or antagonist that alters
 CC the binding of BARD1, B123, BE2, BE14, BE31 or BE445 to BRCA1 or the
 CC biological activity of the BRCA1-BARD1, B123, BE2, BE14, BE31 or BE445
 CC complex. The antibodies can be used to detect BARD1, B123, BE2, BE14,
 CC BE31 or BE445, a specific anti-BARD1 antibody can be used to identify
 CC a patient having or at risk of developing cancer.

XX Sequence 261 AA;

Query Match 62.7%; Score 37; DB 19; Length 261;
 Best Local Similarity 60.0%; Pred. No. 1.1e+02;

Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 2 EAPEPEPIM 11
 DB 219 eaqpeapal 228

RESULT 12

ID AAY90291 standard; Protein; 267 AA.

XX AC AAY90291;
 XX XX

DT	24-OCT-2000	(first entry)
XX		
DE	Human peptidase, HPEP-8 protein sequence.	
KW	Human; peptidase; cell proliferative disorder; arteriosclerosis;	
KW	psoriasis; myelofibrosis; cancer; autoimmune disorder; Crohn's disease;	
KW	inflammatory disorder; AIDS; anaemia; allergy; asthma; arteriosclerosis;	
KW	Grave's disease; multiple sclerosis; scleroderma; infection; diabetes;	
KW	metabolic disorder; Addison's disease; cystic fibrosis; diagnosis;	
KW	glycogen storage disease; obesity; therapy; HPEP-8.	
OS	Homo sapiens.	
PN	WO200042201-A2.	
XX		
PD	20-JUL-2000.	
XX		
PF	11-JAN-2000; 2000WO-US00641.	
XX		
PR	11-JAN-1999; 99US-0172247.	
PR	03-MAY-1999; 99US-0132253.	
PR	27-MAY-1999; 99US-0136653.	
XX		
PA	(INCY-) INCYTE PHARM INC.	
XX		
PI	Bandman O, Hillman JL, Tang YT, Azimzai Y, Baughn MR, Lal P;	
PI	Yue H, Lu DAM;	
XX		
DR	WPI: 2000-482832/42.	
DR	N-PSDB: AAA37664.	
XX		
PT	An isolated polypeptide for diagnosis, prevention and treatment of	
PT	cell proliferative, autoimmune/ inflammatory and metabolic disorders	
PT	comprises a sequence encoding a human peptidase -	
XX		
PS	Claim 2; Page 99; 131pp; English.	
XX		
CC	This sequence represents a human peptidase, designated HPEP-8. The	
CC	invention relates to 18 human peptidases designated HPEP-1 to HPEP-18,	
CC	respectively. The peptidases can be used for treating a disease or	
CC	condition associated with decreased expression or over expression of	
CC	functional human peptidases. The diseases that can be diagnosed,	
CC	prevented and treated include cell proliferative disorders (such as	
CC	arteriosclerosis, psoriasis, myelofibrosis, and cancers),	
CC	autoimmune/inflammatory disorders (such as AIDS, anaemia, allergies,	
CC	Crohn's disease, asthma, arteriosclerosis, Grave's disease, multiple	
CC	sclerosis, and scleroderma), infections, and metabolic disorders (such as	
CC	Addison's disease, diabetes, cystic fibrosis, glycogen storage diseases	
CC	and obesity).	
XX		
XX		
SO	Sequence 267 AA;	
QY	2 EAEPEAEP 9	
	1111111	
Db	247 epepeaep 254	
QY	62.7%; Score 37; DB 21; Length 267;	
	Best Local Similarity 87.5%; Pred. No. 1.2e+02;	
Matches	7; Conservative 0; Mismatches 1; Indels 0; Gaps 0.	
RESULT 13		
AAB20156		
ID	AAB20156 standard; Protein; 267 AA.	
XX		
AC	AAB20156;	
XX		
DT	30-APR-2001 (first entry)	
XX		
DE	Human protein SECP2.	
XX		
XX	SECP2; secreted protein; human; diagnosis; therapy.	

[illegible]

BR 12-MAR-1999; 99US-0124270.
XX
XX (HOMA-) HUMAN GENOME SCI INC.
PA (ROSE/) ROSEN C A.
XX
PI Ruben SM;
XX WPI: 2000-587514/55.
DR N-PSDB; AAF18018.
XX
XX Lung cancer associated gene sequences, referred to as lung cancer
PT antigens, useful for treatment, prevention, and diagnosis of disorders
PT such as lung cancer -
XX
XX Claim 11; Page 961-962; 1425pp; English.
XX
XX Polynucleotide sequences AAF17982 - AAF18424 encode human lung cancer
CC associated proteins represented in AAB58106 - AAB58548. Lung cancer
CC associated proteins and polynucleotide sequences, their agonists, and
CC antagonists may have neuroprotective; cytostatic; cardioactive;
CC immunomodulatory; muscular active general; vulnery; gastrointestinal
CC general; nephrotoxic; antineoplastic; gynecological; or antibacterial
CC activity. The invention also includes antibodies specific for the
CC protein or polynucleotide sequences. The lung cancer associated
CC polynucleotide sequences may be used for detection of lung cancer,
CC chromosome identification, as chromosome markers, and for numerous other
CC diagnostic or research purposes. The proteins may be used to treat
CC disorders such as neural, immune, muscular, reproductive,
CC gastrointestinal, pulmonary, cardiovascular, renal, and proliferative
CC disorders. The proteins may also be used in the treatment of wounds and
CC infectious diseases. Polynucleotide sequences AAF18425 - AAF18433 and
CC peptide AAB58549 are used in the course of the invention for the
CC identification and characterization of the polynucleotide and protein
CC sequences.
XX
XX
SQ Sequence 278 AA:

Query Match 62.7%; Score 37; DB 21; Length 278;
Best Local Similarity 54.5%; Pred. No. 1.2e+02;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
OY 1 HEEPEAEPIIM 11
|||:|:|
DB 57 heakhsrpyim 67

RESULT 15
AAE00235
ID AAE00235 standard; Protein; 302 AA.
XX
XX AAE00235;
AC
XX 13-JUN-2001 (first entry)
DT
XX
XX Protein encoded by Probe 9.
DE
XX
XX Transacylase; taxol; paclitaxel biosynthesis; taxoid; probe;
KW transgenic organism; TAX9.
XX
XX
XX Taxus cuspidata.
OS
XX
XX
XX Key Location/Qualifiers
FH Misc-difference 119
FT /note= "Encoded by TGC"
FT Misc-difference 164
FT /note= "Encoded by ATG"
FT Misc-difference 187
FT /note= "Encoded by TGC"
XX
XX WO200123586-A2.
PN
PD -05-APR-2001.

XX
XX 29-SEP-2000; 2000WO-US27006.
PE
XX
XX 30-SEP-1999; 99US-0411145.
PR 07-DEC-1999; 99US-0457046.
XX
XX
XX (UNIW) UNIV WASHINGTON STATE RES FOUND.
XX
XX
XX Croteau RB, Walker KD, Schoendorf A, Wildung MR;
XX
XX WPI: 2001-245004/25.
DR N-PSDB; AAD03341.
XX
XX
XX New transacylase enzymes, useful for the high yield production of
PT Taxol(TM), related taxoids and useful intermediates in the in the
PT paclitaxel biosynthetic pathway -
XX
XX
XX Claim 1; Page 108-109; 162pp; English.
XX
XX
XX The present sequence is the amino acid sequence of probe 9.
CC Probe 9 is derived from AT-FOR2 and AT-REV1 primers and is
CC used for screening taxus cuspidata TAX9 full length cDNA clone. The
CC probes are useful for the identification of (nucleic acid sequences
CC encoding) transacylases. The probes isolated from the Taxus genus
CC are useful for the synthetic production of Taxol(TM) and related taxoids,
CC as well as intermediates in the paclitaxel biosynthetic pathway. They
CC can also be used for the creation of transgenic organisms that either
CC produce the transacylases for subsequent in vitro use, or produce the
CC transacylases in vivo. The (nucleic acids encoding) transacylases are
CC also useful for creating specific binding agents that recognise the
CC corresponding transacylases. Binding agents include (fragments of)
CC antibodies or any other agent capable of specifically binding to the
CC groups on the proteins.
XX
XX
SQ Sequence 302 AA:

Query Match 62.7%; Score 37; DB 22; Length 302;
Best Local Similarity 70.0%; Pred. No. 1.3e+02;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
OY 2 EAEPEAEPIIM 11
|||:|:|
DB 110 eakpslepim 119

Search completed: January 4, 2002, 08:40:26
Job time: 109 sec

LOCUS A2518927 390 bp DNA GSS 16-OCT-2000
 DEFINITION RPCI-11-67B18. TV RPCI-11 Homo sapiens genomic clone RPCI-11-67B18.
 ACCESSION A2518927
 VERSION A2518927.1 GI:10829921
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 390)
 Zhao, S., Adams, M.D., Nieman, W., Malek, J., de Jong, P. and Venter
 J.C.
 TITLE BAC end sequences of library RPCI-11
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI11-67B18.TJ
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieterdejong.med.bufileo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.bufileo.edu/ordering) or from
 Research Genet cs (info@resgen.com). BAC end search page:
 http://www.tigr.org/cdb/humgen/bac_end_search/bac_end_search.html.
 This BAC end was generated during the Rad process and may have
 higher chance of clone tracking errors.
 Seq primer: T7
 Class: BAC ends.
 FEATURES
 source Location/Qualifiers
 1..390
 /organism="Homo sapiens"
 /db_xref="GDB:7525385"
 /db_xref="taxon:9606"
 /clone_lib="RPCI-11-67B18"
 /clone_lib="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: PBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
 RPCI11 Human Male BAC Library"
 BASE COUNT 59 a 159 c 84 g 87 t 1 others
 ORIGIN
 alignment_scores:
 Quality: 69.00 Length: 12
 Ratio: 6.273 Gaps: 0
 Percent Similarity: 91.667 Percent Identity: 83.333
 alignment_block:
 US-09-444-281-35 x A2518927/rev ..
 Align seg 1/1 to reverse of: A2518927 from: 1 to: 390
 2 LeuLysLysTrpProTrpTrpProTrpPAAGAGLys 13
 |||||:|||| |||||:|||||||:|||||:|||||
 189 TTACAAAATAATCCCTGCTGGCCCTGAGAGAGAG 154
 seq_name: gb_est1:BE237369
 seq_documentation_block:
 LOCUS BE237369 415 bp mRNA EST 25-APR-2001
 DEFINITION 146629 MARC 4BOV Bos taurus cDNA 5', mRNA sequence.
 ACCESSION BE237369
 VERSION BE237369.1 GI:9022087
 KEYWORDS EST.
 SOURCE COW.
 ORGANISM Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 Bovidae; Bovinae; Bos.
 1 (bases 1 to 415)
 Smith, T.P.L., Grosse, W.M., Freking, B.A., Roberts, A.J., Stone, R.T.,
 Casas, E., Wray, J.E., White, J., Cho, J., Fahrenkrug, S.C., Bennett,
 G.L., Heaton, M.P., Laegreid, W.W., Rohrer, G.A., Chitko-McKown, C.G.,
 Perlea, G., Holt, L., Karameycheva, S., Liang, F., Quackenbush, J. and
 Keeler, J.W.
 TITLE Sequence evaluation of four pooled-tissue normalized bovine cDNA
 JOURNAL libraries and construction of a gene index for cattle
 MEDLINE Genome Res. 11 (4), 626-630 (2001)
 COMMENT 2180013
 Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smith@email.marc.usda.gov
 Single pass sequencing. Bases called and alt-trimmed with phred
 v0.980904.e. Vector identified by cross-match with the -minscore 18
 and -mismatch 12 options.
 PCR Primers
 FORWARD: AGGAACACGCTATGACCAT
 BACKWARD: GTTTCACGCTACGACG
 Plate: 47 row: J column: 10
 Seq primer: ATTATGGTGACATATAC.
 FEATURES
 source Location/Qualifiers
 1..415
 /organism="Bos taurus"
 /db_xref="taxon:9913"
 /clone_lib="MARC 4BOV"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /note="Vector: pCMV SPORT6; Site_1: XbaI; Site_2: XhoI;
 Library made from pooled tissue from day 20 and day 40
 embryos."
 BASE COUNT 106 a 97 c 120 g 91 t 1 others
 ORIGIN
 alignment_scores:
 Quality: 68.00 Length: 8
 Ratio: 8.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000
 alignment_block:
 US-09-444-281-35 x BE237369 ..
 Align seg 1/1 to: BE237369 from: 1 to: 415
 4 LysTrpProTrpTrpProTrpPArg 11
 |||||:|||||:|||||:|||||:|||||
 280 AATGGCCATGCTGGCCCTTGGCGC 303
 seq_name: gb_est1:A0089922
 seq_documentation_block:
 LOCUS A0089922 446 bp mRNA EST 19-APR-2000
 DEFINITION A0089922 Hordeum vulgare subsp. vulgare upper three leaves at
 heading stage Hordeum vulgare subsp. vulgare cDNA clone
 haruna_lib1_121, mRNA sequence.
 ACCESSION A0089922
 VERSION A0089922.1 GI:7613350
 KEYWORDS EST.
 SOURCE Hordeum vulgare subsp. vulgare.
 ORGANISM Hordeum vulgare subsp. vulgare.
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Poideae
 ; Triticeae; Hordeum.
 1 (bases 1 to 446)
 Sato, K., Takahashi, H. and Takeda, K.
 TITLE Hordeum vulgare subsp. vulgare cDNA clone
 JOURNAL Unpublished (2000)

COMMENT

Contact: Kazuhiro Sato
Research Institute for Bioresources
Okayama University, Barley Germplasm Center
Chuo 2-20-1 Kurashiki, Okayama 710-0046, Japan
Email: kase@oerib.okayama-u.ac.jp,
URL: http://www.rib.okayama-u.ac.jp/barley/
location/Qualifiers

FEATURES

source
1. 446
/organism="Hordeum vulgare subsp. vulgare"
/cultivar="Haruna Nijo"
/db_xref="taxon:112509"
/clone="haruna_1lib1_121"
/clone_1lb="Hordeum vulgare subsp. vulgare Upper three
leaves at heading stage"
/tissue_type="Upper three leaves at heading stage"
BASE COUNT 89 a 130 c 149 g 76 t 2 others
ORIGIN

alignment_scores:

Quality: 68.00 Length: 8
Ratio: 8.500 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-35 x AU089922/rev ..

Align seg 1/1 to reverse of: AU089922 from: 1 to: 446

5 TrpProtTrpProtTrpParGarg 12
|||||
188 TGGCGCTGGCTGGCGCGCGCA 165

seq_name: gb_est1:AU198144

seq_documentation_block:

LOCUS AU198144 446 bp mRNA EST 12-JUL-2001
DEFINITION AU198144 Rice green shoot Oryza sativa cDNA clone S15951, mRNA
sequence.
ACCESSION AU198144
VERSION AU198144.1 GI:14714211
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzaceae; Oryza.
1 (bases 1 to 446)
Sasaki, T. and Yamamoto, K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@dr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16019_97A.

REFERENCE

AUTHORS
TITLE
JOURNAL
COMMENT
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@dr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S15951_97A.

FEATURES

source
1. 446
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="S15951"
/clone_1lb="Rice green shoot"
/note="Green shoot (8 days old)"
BASE COUNT 82 a 153 c 161 g 50 t
ORIGIN

alignment_scores:

Quality: 68.00 Length: 11
Ratio: 6.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-09-444-281-35 x AU198144/rev ..

Align seg 1/1 to reverse of: AU198144 from: 1 to: 446

3 LysLysTrpProtTrpProtTrpParGargLys 13
:::|||||
351 CGCGCTGGCTGGCGCGCGCGCGG 319

seq_name: gb_est1:AU198162

seq_documentation_block:

LOCUS AU198162 448 bp mRNA EST 12-JUL-2001
DEFINITION AU198162 Rice green shoot Oryza sativa cDNA clone S16019, mRNA
sequence.
ACCESSION AU198162
VERSION AU198162.1 GI:14714231
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehrhartoideae; Oryzaceae; Oryza.
1 (bases 1 to 448)
Sasaki, T. and Yamamoto, K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@dr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16019_97A.

REFERENCE

AUTHORS
TITLE
JOURNAL
COMMENT
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@dr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16019_97A.

FEATURES

source
1. 448
/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="S16019"
/clone_1lb="Rice green shoot"
/note="Green shoot (8 days old)"
BASE COUNT 85 a 146 c 160 g 57 t
ORIGIN

alignment_scores:

Quality: 68.00 Length: 11
Ratio: 6.800 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-09-444-281-35 x AU198162/rev ..

Align seg 1/1 to reverse of: AU198162 from: 1 to: 448

3 LysLysTrpProtTrpProtTrpParGargLys 13
:::|||||
314 CGCGCTGGCTGGCGCGCGCGCGG 282

seq_name: gb_est1:AU089934

seq_documentation_block:

LOCUS AU089934 475 bp mRNA EST 19-APR-2000
DEFINITION AU089934 Hordeum vulgare subsp. vulgare upper three leaves at
heading stage Hordeum vulgare subsp. vulgare cDNA clone
haruna_1lib1_134, mRNA sequence.

```

ACCESSION      AU089934
VERSION        AU089934.1
KEYWORDS       EST
SOURCE         Hordeum vulgare subsp. vulgare.
ORGANISM       Hordeum vulgare subsp. vulgare
REFERENCE      Sato, K., Takahashi, H. and Takeda, K.
AUTHORS        1 (bases 1 to 475)
TITLE          Hordeum vulgare subsp. vulgare cDNA clone
JOURNAL        Unpublished (2000)
COMMENT        Contact: Kazuhiko Sato
                Research Institute for Bioreources
                Okayama University, Barley Germplasm Center
                Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
                Email: kazsato@rib.okayama-u.ac.jp/,
                URL:http://www.rib.okayama-u.ac.jp/barley/.
FEATURES       location/Qualifiers
               1..475
               /organism="Hordeum vulgare subsp. vulgare"
               /cultivar="Hartuna Nijo"
               /db_xref="taxon:112509"
               /clone="hartuna libl 134"
               /clone_lib="Hordeum vulgare subsp. vulgare Upper three
               leaves at heading stage"
               /tissue_type="Upper three leaves at heading stage"
BASE COUNT     100 a 137 c 155 g 83 t
ORIGIN
alignment_scores:
    Quality:   68.00           Length:   8
    Ratio:     8.500           Gaps:     0
Percent Similarity: 100.000 Percent Identity: 100.000
alignment_block:
US-09-444-281-35 x AU089934/rev ..
Align seg 1/1 to reverse of: AU089934 from: 1 to: 475
5 TrpproTPTPrpTrprArGy 12
|||||
194 TGCCCGTGCTGGCCGTGGCGGC GA 171
seq_name: gb_est1:AU082117
seq_documentation_block:
LOCUS          AU082117             578 bp            mRNA            EST            04-FEB-2000
DEFINITION    AU082117 Rice panicle at ripening stage Oryza sativa cDNA clone
EIL1611, mRNA sequence.
ACCESSION     AU082117
VERSION       AU082117.1 GI:6727452
KEYWORDS      EST.
SOURCE        Oryza sativa.
ORGANISM      Oryza sativa
EXTRACT       Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
                Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
                Ehrhartoideae; Oryzaceae; Oryza.
1 (bases 1 to 578)
I (bases 1 to 578)
Sasaki,T. and Yamamoto,K.
Rice cDNA from panicle at ripening stage (2000)
Unpublished (2000)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasakieabr.affrc.go.jp, URL:http://rpg.dna.affrc.go.jp/
PROJECT       "RGP".
FEATURES       Location/Qualifiers
               1..578

```

```

/organism="Oryza sativa"
/strain="Nipponbare"
/db_xref="taxon:4530"
/clone="E1611"
/clone_lib="Rice panicle at ripening stage"
/dev_stage="ripening stage"
/note="Organ: panicle; Rice cDNA from panicle at ripening stage"
BASE COUNT      129 a      180 c      179 g      90 t
ORIGIN

alignment_scores:
  Quality:      68.00      Length:      11
  Ratio:        6.800      Gaps:      0
Percent Similarity: 90.909      Percent Identity: 63.636

alignment_block:
US-09-444-281-35 x AU082117/rev      ..

Align seg 1/1 to reverse of: AU082117 from: 1 to: 578

3 LysLysTTPProTTPProTTPProTPArGArGlys 13
:::|||||
348 CGCCGCTGCTGCTGCTGCTGCTGCGCGGCGG 316

seq_name: gb_est1:AU198258

seq_documentation_block:
LOCUS      AU198258      352 bp      mRNA      EST      12-JUL-2001
DEFINITION AU198258 Rice green shoot Oryza sativa cDNA clone S16389, mRNA
sequence.
ACCESSION  AU198258
VERSION    AU198258.1 GI:14714335
KEYWORDS   EST.
SOURCE     Oryza sativa.
ORGANISM   Oryza sativa
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Erihartoideae; Oryzaceae; Oryza.
1 (bases 1 to 352)
Sasaki,T. and Yamamoto,K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@agr.affrc.go.jp, URL:http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16389_96Z.

FEATURES
    source
        1..352
            location/Qualifiers
                organism="Oryza sativa"
                strain="Nipponbare"
                db_xref="taxon:4530"
                clone="S16389"
                clone_lib="Rice green shoot"
                note="Green shoot (8 days old)"
BASE COUNT      95 a      95 c      104 g      51 t      7 others
ORIGIN

alignment_scores:
  Quality:      66.00      Length:      10
  Ratio:        7.333      Gaps:      0
Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
US-09-444-281-35 x AU198258/rev      ..

```

Align seg 1/1 to reverse of: AU198258 from: 1 to: 352

3 Lyslystiprrotrpprpararg 12
 170 CGCGCTGGCTTGTCGACGCG 141

seq_name: gb_est1:BE453064

seq_documentation_block:

LOCUS BE453064 434 bp mRNA EST 26-JUL-2000
 DEFINITION 894068B05.x1 C. reinhardtii CC-1690, normalized, lambda Zap II
 Chlamydomonas reinhardtii cDNA, mRNA sequence.

ACCESSION BE453064
 KEYWORDS BE453064.1 GI:9460101
 EST.

SOURCE Chlamydomonas reinhardtii.
 ORGANISM Chlamydomonas reinhardtii
 Eukaryota: Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
 Chlamydomonadaceae; Chlamydomonas.

REFERENCE 1 (bases 1 to 434)

AUTHORS Grossman, A., Davies, J., Federspiel, N., Harris, E., Lefebvre, P.,
 McDermott, J. P., Sillitow, C., Stern, D. and Surzycki, R.

TITLE Analyses of the Chlamydomonas reinhardtii Genome: A Model,
 Unicellular System for Analyzing Gene Function and Regulation in
 Vascular Plants; project phase 2

JOURNAL Unpublished (2000)
 COMMENT Contact: Elizabeth H. Harris
 DCMB Box 91000
 Duke University
 Durham, NC 27708-1000, USA
 Tel: 919 613 8164
 Fax: 919 613 8177
 Email: chlamy@duke.edu.

FEATURES
 source Location/Qualifiers

1..434
 /organism="Chlamydomonas reinhardtii"
 /strain="CC-1690 wild type mt+ 21gr"
 /db_xref="taxon:3055"
 /clone_lib="C. reinhardtii CC-1690, normalized, lambda Zap
 II"

/note="Vector: Bluescript II SK-; Site_1: EcoRI; Site_2:
 XhoI; This library, constructed by John Davies and Jeffrey
 McDermott, combines cDNAs from CC-1690 cells grown to
 mid-log phase in TAP (acetate-containing) medium in the
 light, TAP medium in the dark, HS (minimal) medium in
 ambient levels of CO2 and HS medium bubbled with 5% CO2.
 PolyA mRNA was purified from each sample, pooled and cDNA
 synthesized. The cDNA was directionally cloned into lambda
 Zap II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
 Bluescript II SK- plasmids were excised from the lambda
 Zap clones by superinfection with ExAssist (Stratagene)
 phage. The library was normalized using method 4 described
 in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT 110 a 146 c 86 g 92 t
 ORIGIN

alignment_scores:
 Quality: 66.00 Length: 11
 Ratio: 7.333 Gaps: 0
 Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:
 US-09-444-281-35 x BE453064/rev ..

Align seg 1/1 to reverse of: BE453064 from: 1 to: 434

1 ltleulyslystiprrotrpprpararg 11
 403 AATAATGTCGATGCGCATGTCGACGCG 371
 seq_name: gb_est2:BG305131

seq_documentation_block:

LOCUS BG305131 445 bp mRNA EST 22-FEB-2001
 DEFINITION F195a09.y1 Zebrafish adult retina cDNA Danio rerio cDNA clone
 414409.5, mRNA sequence.

ACCESSION BG305131
 KEYWORDS BG305131.1 GI:13102658
 EST.

SOURCE zebrafish.
 ORGANISM Danio rerio
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Actinopterygii; Neopterygii; Teleostei; Euteleostei; Ostariophysi;
 Cypriniformes; Cyprinidae; Rasbora; Danio.

REFERENCE 1 (bases 1 to 445)

AUTHORS Clark, M., Johnson, S. L., Lehnach, H., Lee, R., Li, F., Marra, M., Eddy,
 S., Hillier, L., Kucaba, T., Martin, J., Beck, C., Wylie, T., Underwood,
 K., Stepien, M., Theising, B., Allen, M., Bowers, Y., Person, B.,
 Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter, E.,
 Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R.
 and Wilson, R.

TITLE Washu zebrafish EST Project 1998

JOURNAL Unpublished (1998)
 COMMENT Contact: Stephen L. Johnson
 Washington University School of Medicine
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
 Tel: 314 286 1800
 Fax: 314 286 1810

Email: zbrfish@watson.wustl.edu
 Library constructed by: Susan E. Brockerhoff DNA Sequencing by:
 Washington University Genome Sequencing Center Clone distribution:
 Ressourcenzentrumprimatardenbank, Berlin, Germany (web address:
 www.rzpd.de)

Seq primer: T3 EF from Amersham
 High quality sequence stop: 419.

FEATURES
 source Location/Qualifiers

1..445
 /organism="Danio rerio"
 /strain="wild-type"
 /db_xref="taxon:7955"
 /clone_lib="414409"
 /clone_lib="Zebrafish adult retina cDNA"
 /sex="mixed"
 /dev_stage="1-2 years"
 /lab_host="E. Coli XL1-Blue MRF' (XL1-Blue MRF')"
 /note="Vector: lambda Zap II (Bluescript SK-); Site_1:
 EcoRI; Site_2: SalI; This Zebrafish library was
 constructed by Dr. Susan E. Brockerhoff (email:
 sbrocker@u.washington.edu) RZPD library number: 760"

BASE COUNT 97 a 140 c 119 g 89 t
 ORIGIN

alignment_scores:
 Quality: 66.00 Length: 11
 Ratio: 7.333 Gaps: 0
 Percent Similarity: 81.818 Percent Identity: 72.727

alignment_block:

US-09-444-281-35 x BG305131/rev ..

Align seg 1/1 to reverse of: BG305131 from: 1 to: 445

3 Lyslystiprrotrpprpararglys 13

252 AAACGTGGCCCTGTCGACGACGAGAG 220

seq_name: gb_gss:AZ368740

seq_documentation_block:

LOCUS AZ368740 600 bp DNA GSS 02-OCT-2000
 DEFINITION 1M0118021R Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0118021 R, DNA sequence.

ACCESSION AZ368740
 VERSION AZ368740.1 GI:10482440

KEYWORDS
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE
AUTHORS 1 (bases 1 to 600)
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
Islam,H., Longacre,S., Mahmood,M., Meenen,E., Pedersen,T., Reilly
M., Rose,M., Rose,R., Stokes,R., Tinney,A., von Niederhausen,A.
and Wright,D., Weiss,R.
Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts
JOURNAL
COMMENT
TITLE
Plasmid inserts
Contact: Robert B. Weiss
University of Utah Genome Center
University of Utah
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA
Tel: 801 585 5606
Fax: 801 585 7177
Email: ddunn@genetics.utah.edu
Insert length: 10000 Std Error: 0.00
Plate: 0118 Row: 0 Column: 21
Seq primer: CACACAGGAACACGCTATGACC
Class: plasmid ends
High quality sequence stop: 600.
Location/Qualifiers
1. 600
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UUCG1M0118021"
/clone_lib="Mouse 10kb plasmid UUCG1M library"
/sex="Male"
/lab_host="E. coli strain XL10-Gold, T1-resistant, F-"
/note="Vector: pMD22ny; Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydronically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The
adaptor DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of pMD22 (g11473211419b/AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adaptor mouse DNA was annealed to
adaptor vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

BASE COUNT 173 a 108 c 181 g 138 t
ORIGIN

alignment_scores:
Quality: 60.00 Length: 9
Ratio: 7.333 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-09-444-281-35 x A2368740/rev ..
Align seg 1/1 to reverse of: A2368740 from: 1 to: 600

2 LysLysLysTrpProTrpTrpProTrp 10
|||||:|||||:|||||:|||||:|||||
384 TTACGGAGGTGGCCATGCTGCTGG 358

seq_name: gb_est2:BE977776

seq documentation block:
LOCUS BE977776 255 bp mRNA EST 04-OCT-2000
DEFINITION bs68h10.y1 Drosophila melanogaster adult testis library Drosophila
melanogaster cDNA clone bs68h10 5', mRNA sequence.
ACCESSION BE977776
VERSION BE977776.1 GI:10608588
KEYWORDS EST.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 255)
REFERENCE Andrews, J., Bouffard, G. and Oliver, B.
Drosophila melanogaster testis expressed sequence tags
JOURNAL
COMMENT
TITLE
Drosophila melanogaster testis expressed sequence tags
Contact: Brian Oliver
Laboratory of Cellular and Developmental Biology
NIDDK, National Institutes of Health
6 Center Drive MSC 2715, Bldg 6, Rm B1-13, Bethesda, MD 20892 USA
Fax: (301) 496 5239
Email: oliver@helix.nih.gov
http://www.nidk.nih.gov/intram/people/boliver.htm
Tissue isolation and library construction performed at the National
Institute of Diabetes and Digestive and Kidney Diseases, NIH (see
http://www.nidk.nih.gov/intram/people/boliver.htm). DNA sequencing
and analyses performed by National Institutes of Health Intramural
Sequencing Center (NISC; see http://www.nisc.nih.gov).
Plate: 68 Row: h Column: 10
Seq primer: MJ3Rpl reverse primer (ABI).
Location/Qualifiers
1. 255
/organism="Drosophila melanogaster"
/strain="y1 w[67c1]/Y"
/db_xref="taxon:7227"
/clone="bs68h10"
/clone_lib="Drosophila melanogaster adult testis library"
/sex="male"
/dev_stage="1-5 day adult"
/lab_host="SOLR (Stratagene)"
/note="Organ: testis; Vector: pBluescript SK (Stratagene);
Site_1: Bcor 1; Site_2: Xho 1; Testes dissected from 1-5
day adult y1 w[67c1]/Y males raised at 25°C. RNA
isolated using Trizol (Life Technologies) and a single
round of Poly(A)+ selection using Oligotex (Qiagen). cDNA
library constructed using Stratagene ZAP-cDNA synthesis
kit. Oligo dT-primed, size fractionated -1-6 kb, and
directionally cloned at EcoRI and XhoI in Uni-ZAP XR.
Following a single round of amplification pBluescript SK
phagemids were mass excised. A distribution channel for
clones is being sought, but not currently available.
Requests for clones cannot be honored."

BASE COUNT 47 a 60 c 101 g 47 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 15
Ratio: 5.909 Gaps: 1
Percent Similarity: 73.333 Percent Identity: 60.000

alignment_block:
US-09-444-281-35 x BE977776 ..
Align seg 1/1 to: BE977776 from: 1 to: 255

3 LysLysTrpPro.....TrpTrpProTrpArgLys 13
|||||:|||||:|||||:|||||:|||||
185 AAGAGCTGCCCCGTGAGGAAAAGTGTGCTGCTGGAGGAGMA 229

seq_name: gb_est1:AV641634

```

seq_documentation_block: 371 bp, mRNA, EST, 15-DEC-2000
LOCUS AV641634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
DEFINITION AV641634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
CDNA clone HCL037h03_r 5' mRNA sequence.
ACCESSION AV641634
VERSION AV641634.1 GI:10784962
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii.
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.
REFERENCE 1 (bases 1 to 371)
AUTHORS Asanizu,E., Miura,K., Kuchio,K., Inoue,Y., Fukuzawa,H., Ohnaya,K.,
Nakamura,Y. and Tabata,S.
TITLE Generation of expressed sequence tags from low-CO2 and high-CO2
adapted cells of Chlamydomonas reinhardtii
JOURNAL DNA Res. 7 (5), 305-307 (2000)
MEDLINE 20539644
COMMENT Contact: Erika Asanizu
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asanizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.
Location/Qualifiers
1. 371
/organism="Chlamydomonas reinhardtii"
/strain="C9"
/db_xref="taxon:3055"
/clone="HCL037h03_r"
/clone_lib="Chlamydomonas reinhardtii 5% CO2"
/note="Vector: pBluescriptII SK-; Site:1: EcoRI; Site:2:
XhoI. The cDNA library was constructed from cells cultured
in a medium with bubbling air containing 5% carbon
dioxide"
BASE COUNT 73 a 106 c 141 g 51 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10
Ratio: 7.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-35 x AV641634 ...
Align seg 1/1 to: AV641634 from: 1 to: 371

3 LysLysTrpProTrpTrpProTrpArgArg 12
+++++|||||
241 GAACGGTGGCGGTGGTGGCGCGCGG 270

seq_name: gb_est1:BE129188

seq_documentation_block: 386 bp, mRNA, EST, 21-JUN-2000
LOCUS BE129188 Chlamydomonas reinhardtii CC-1690, normalized, Lambda Zap II
DEFINITION 894021E12.Y1 C. reinhardtii CC-1690, normalized, Lambda Zap II
Chlamydomonas reinhardtii cDNA, mRNA sequence.
ACCESSION BE129188
VERSION BE129188.1 GI:8576551
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii.
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.
REFERENCE 1 (bases 1 to 386)
AUTHORS Grossman,A., Davies,J., Federspiel,N., Harris,E., Lefebvre,P.,
McDermott,J.P., Silflow,C., Stern,D. and Surzycki,R.
TITLE Analyses of the Chlamydomonas reinhardtii Genome: A Model,
Unicellular System for Analyzing Gene Function and Regulation in
Vascular Plants; project phase 2
JOURNAL Unpublished (2000)
COMMENT Contact: Elizabeth H. Harris

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```

DCMB Box 91000
Duke University
Durham, NC 27708-1000, USA
Tel: 919 613 8164
Fax: 919 613 8177
Email: chlamydeduke.edu.
Location/Qualifiers
1. 386
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type ml+ 21gr"
/db_xref="taxon:3055"
/clone_lib="C. reinhardtii CC-1690, normalized, Lambda Zap
II"
/note="Vector: pBluescript II SK-; Site:1: EcoRI; Site:2:
XhoI. This library, constructed by John Davies and Jeffrey
McDermott, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP (acetate-containing) medium in the
light, TAP medium in the dark, HS (minimal) medium in
ambient levels of CO2 and HS medium bubbled with 5% CO2.
PolyA mRNA was purified from each sample, pooled and cDNA
synthesized. The cDNA was directionally cloned into lambda
Zap II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
pBluescript II SK- plasmids were excised from the lambda
Zap clones by superinfection with ExAssist (Stratagene)
phage. The library was normalized using method 4 described
in Bonaldo et al (1996) Genome Research 6: 791-806."
BASE COUNT 79 a 109 c 142 g 56 t
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10
Ratio: 7.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-09-444-281-35 x BE129188 ..
Align seg 1/1 to: BE129188 from: 1 to: 386

3 LysLysTrpProTrpTrpProTrpArgArg 12
+++++|||||
246 GAACGGTGGCGGTGGTGGCGCGCGG 275

```

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:23 : Search time 50.17 seconds
(without alignments)
37.902 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKKPMWPMWRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues
Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: SPTEMBL_17:*
2: sp.archaea:*
3: sp.bacteria:*
4: sp.fungi:*
5: sp.human:*
6: sp.invertebrate:*
7: sp.mammal:*
8: sp.mhc:*
9: sp.phage:*
10: sp.plant:*
11: sp rodent:*
12: sp.virus:*
13: sp.vertebrate:*
14: sp.unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	57	62.6	723	12	09DUC4
2	54	59.3	1173	12	0990M4
3	54	59.3	1173	12	0990M3
4	54	59.3	1173	12	0990M2
5	54	59.3	1173	12	0990M1
6	54	59.3	1173	12	0990M1
7	52	57.1	746	12	094712
8	52	57.1	746	12	094712
9	51	56.0	399	4	09YANI
10	50	54.9	327	10	09AUN3
11	49	53.8	148	5	026590
12	49	53.8	298	1	09Y806
13	49	53.8	467	5	019573
14	49	53.8	528	5	026589
15	49	53.8	528	5	09Y57
16	49	53.8	725	12	09DUC9
17	49	53.8	802	5	096398
18	49	53.8	1245	3	09Y7V5
19	49	53.8	1940	5	002456

20	48	52.7	49	12	09DTR80	09dtr80 tt virus. o
21	48	52.7	111	5	018753	018753 caenorhabdi
22	48	52.7	428	11	09JMG0	09jmg0 mus musculu
23	48	52.7	431	11	099ML4	099ml4 mus musculu
24	48	52.7	748	12	09DTR81	09dtr81 tt virus. o
25	47.5	52.2	114	2	09XBC2	09xbc2 streptomyce
26	47	51.6	141	11	09CZAI	09czai mus musculu
27	47	51.6	165	10	09SNN3	09snn3 oryza sativ
28	47	51.6	504	2	P96143	P96143 thermoactin
29	46.5	51.1	352	2	P73417	P73417 synchocyst
30	46.5	51.1	620	12	091H07	091h07 avian infec
31	46.5	51.1	621	12	066196	066196 avian infec
32	46.5	51.1	621	12	091H14	091h14 avian infec
33	46.5	51.1	621	12	091H13	091h13 avian infec
34	46.5	51.1	621	12	091H11	091h11 avian infec
35	46.5	51.1	621	12	091H10	091h10 avian infec
36	46.5	51.1	621	12	091H15	091h15 avian infec
37	46.5	51.1	621	12	091H12	091h12 avian infec
38	46.5	51.1	621	12	091H09	091h09 avian infec
39	46.5	51.1	621	12	091H08	091h08 avian infec
40	46.5	51.1	621	12	0991L8	0991l8 avian infec
41	46.5	51.1	621	12	0991L7	0991l7 avian infec
42	46.5	51.1	621	12	0991L6	0991l6 avian infec
43	46.5	51.1	625	12	09QCP6	09qcp6 avian infec
44	46.5	51.1	630	12	066197	066197 avian infec
45	46.5	51.1	630	12	098WQ0	098wq0 avian infec

ALIGNMENTS

RESULT 1
ID 09DUC4 PRELIMINARY; PRT; 723 AA.
AC 09DUC4;
DT 01-MAR-2001 (TEMBLrel. 16, Created)
DT 01-MAR-2001 (TEMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TEMBLrel. 17, Last annotation update)
DE ORF1.
OS TTR virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RA Okamoto H.;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RX PubMed-11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TTV viruses in humans and nonhuman primates and their
RL phylogenetic relatedness.";
RL Virology 277:368-378(2000).
DR EMBL: AB041959; BAB19313.1;
DR InterPro: IPR001563; Serine_carpept.
DR PROSITE: PS00131; CARBOXYPEPT_SER_SER; UNKNOWN_1.
SO SEQUENCE 723 AA; 85393 MW; 232D003098766344 CRC64;

Query Match 62.68; Score 57; DB 12; Length 723;
Best Local Similarity 100.0%; Pred. No. 4.1;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 6 PWMPPRR 12
| | | | |
Db 2 PWMPPRR 8

RESULT 2
0990M4

ID 0990M4 PRELIMINARY; PRT; 1173 AA.
AC 0990M4;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344186; AAK32188.1;
SQ SEQUENCE 1173 AA; 128669 MW; ABC6E0A75EBDBA4 CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
DB 1113 KWPMPW 1119

RESULT 3
0990M3 PRELIMINARY; PRT; 1173 AA.
AC 0990M3;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344187; AAK32189.1;
SQ SEQUENCE 1173 AA; 128683 MW; 9E236816082A81A CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
DB 1113 KWPMPW 1119

RESULT 4
0990M2 PRELIMINARY; PRT; 1173 AA.
AC 0990M2;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.

OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344188; AAK32190.1;
SQ SEQUENCE 1173 AA; 128653 MW; 8B658FCBBD1842DA CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
DB 1113 KWPMPW 1119

RESULT 5
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AC 0990M1;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344189; AAK32191.1;
SQ SEQUENCE 1173 AA; 128760 MW; B73A165A6270152A CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1173;
Best Local Similarity 85.7%; Pred. No. 16;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KWPMPW 10
DB 1113 KWPMPW 1119

RESULT 6
084712 PRELIMINARY; PRT; 1383 AA.
AC 084712;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DE SPIKE PROTEIN.
OS Porcine epidemic diarrhea virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=28295;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=94231173; PubMed=8176382;

RA Duarte M., Laude H.;
RT "Sequence of the spike protein of the porcine epidemic diarrhoea virus."
RL J. Gen. Virol. 75:1195-1200(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-BRI/87;
RX MEDLINE=93389433; PubMed=8397280;
RA Bridgen A., Duarte M., Tobler K., Laude H., Ackermann M.;
RT "Sequence determination of the nucleocapsid protein gene of the porcine epidemic diarrhoea virus confirms that this virus is a transmissible gastroenteritis virus."
RL J. Gen. Virol. 74:1795-1804(1993).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-BRI/87;
RX MEDLINE=94120721; PubMed=8291230;
RA Duarte M., Tobler K., Bridgen A., Rasschaert D., Ackermann M., Laude H.;
RT "Sequence analysis of the porcine epidemic diarrhoea virus genome between the nucleocapsid and spike protein genes reveals a polymorphic ORF."
RL Virology 198:466-476(1994).
DR EMBL: Z25483; CAA80971.1; -;
DR InterPro: IPR002551; Corona_S1.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
RT CONFLICT 422 422 Y -> N (IN REF. 1).
SQ SEQUENCE 1383 AA; 151404 MW; 741C84D5DD3BDCAD CRC64;

Query Match 59.3%; Score 54; DB 12; Length 1383;
Best Local Similarity 85.7%; Pred. No. 18;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 4 KMPMPW 10
Db 1322 KMPMPW 1328

RESULT 7
Q9JH31
ID 09JH31 PRELIMINARY; PRT; 746 AA.
AC 09JH31;
DT 01-OCT-2000 (TEMBLrel. 15, Created)
DT 01-OCT-2000 (TEMBLrel. 15, Last sequence update)
DE ORF1.
OS Virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-TJN02;
RA Okamoto H.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-TJN02;
RA Ukita M., Okamoto H., Nishizawa T., Tawara A., Takahashi M., Iizuka H., Miyakawa Y., Mayumi M.;
RT "The entire nucleotide sequences of two distinct TT virus (TTV) isolates (TJN01 and TJN02) remotely related to the original TTV isolates."
RL Arch. Virol. 0:0-0(2000).
DR EMBL: AB028669; BAA94878.1; -;
SQ SEQUENCE 746 AA; 88561 MW; E0B22953AE764E3E CRC64;

Query Match 57.1%; Score 52; DB 12; Length 746;
Best Local Similarity 66.7%; Pred. No. 19;

Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
Qy 5 WPMWPRK 13
Db 3 WPMWPRR 11

RESULT 8
Q9HKX3
ID 09HKX3 PRELIMINARY; PRT; 1018 AA.
AC 09HKX3;
DT 01-MAR-2001 (TEMBLrel. 16, Created)
DT 01-MAR-2001 (TEMBLrel. 16, Last sequence update)
DT 01-JUN-2001 (TEMBLrel. 17, Last annotation update)
DE CONSERVED HYPOTHETICAL MEMBRANE PROTEIN.
GN TAO470.
OS Thermoplasma acidophilum.
OC Archaea; Euryarchaeota; Thermoplasmatales; Thermoplasmaceae;
OC Thermoplasma.
OX NCBI_TaxID=2303;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-DSM 1728;
RX MEDLINE=20479972; PubMed=11029001;
RA Ruepp A., Graml W., Santos-Martinez M.-L., Koretke K.K., Volker C., Mewes H.-W., Fishman D., Stocker S., Lupas A.N., Baumeister W.;
RT "The genome sequence of the thermophilic scavenger Thermoplasma acidophilum."
RL Nature 407:508-513(2000).
DR EMBL: AL445064; CAC11612.1; -;
DR InterPro: IPR000731; HMGCR_Patched_5TM.
DR PROSITE: PS50156; SSD; 1.
KW Complete proteome.
SQ SEQUENCE 1018 AA; 112323 MW; 83EE84D3C74B852 CRC64;

Query Match 57.1%; Score 52; DB 1; Length 1018;
Best Local Similarity 66.7%; Pred. No. 25;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
Qy 1 ILKMPWP 9
Db 1004 ILKMPWP 1012

RESULT 9
Q9Y4N1
ID 09Y4N1 PRELIMINARY; PRT; 299 AA.
AC 09Y4N1;
DT 01-NOV-1999 (TEMBLrel. 12, Created)
DT 01-MAY-2000 (TEMBLrel. 13, Last sequence update)
DT 01-MAY-2000 (TEMBLrel. 13, Last annotation update)
DE HYPOTHETICAL 34.0 KDA PROTEIN (FRAGMENT).
GN DKFZP434C192.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=TESTIS;
RA Ansoorge W., Winkler U., Mewes H.W., Gassenhuber J., Wiemann S.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL096753; CAB46428.2; -;
KW Hypothetical protein.
FT NON_TER 1
SQ SEQUENCE 299 AA; 34032 MW; 6B8DB60E6A88239A CRC64;

Query Match 56.0%; Score 51; DB 4; Length 299;
Best Local Similarity 85.7%; Pred. No. 11;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 6 PWMWR 12
 ||||| 1
 Db 37 PWMWR 43

RESULT 10
 ID 09AUN3 PRELIMINARY; PRT; 327 AA.
 AC 09AUN3:
 DT 01-JUN-2001 (TREMBLrel. 17, Created)
 DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
 DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE HYPOTHEITICAL PROTEIN.
 OS Oryza sativa (Rice).
 CC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 CC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 CC Eriactoidae; Oryzae; Oryza.
 NCBI_TaxID=4530;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Spiegel L.A., Kirchoff K.A., de la Bastide M., Preston R.R.,
 RA Nascimiento L.U., Vil M.D., Baker J.P., Miller B., Cunius D.M.,
 RA Kuit K.H., Rodriguez S., Santos L., Zutavern T., Ballja V.S.,
 RA Shah R.S., Bahret A., Bal H.P., O'Shaughnessy A., Dedila N.N.,
 RA McCombie W.R.;
 RT "Genomic Sequence For Oryza sativa, Nipponbare Strain, Chromosome X,
 RT Clone OSUBA0058B19, Complete Sequence."
 RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AC083945; AAK13143.1;
 SQ SEQUENCE 327 AA; 36672 MW; 5CCA9080664BD0CA CRC64;

Query Match 54.9%; Score 50; DB 10; Length 327;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 7 WWPWR 12
 ||||| 1
 Db 119 WWPWR 124

RESULT 11
 ID 026590 PRELIMINARY; PRT; 148 AA.
 AC 026590:
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-OCT-2000 (TREMBLrel. 15, Last annotation update)
 DE MYOSIN HEAVY CHAIN (FRAGMENT).
 OS Schistosoma mansoni (Blood fluke).
 CC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
 CC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
 CC Schistosoma
 NCBI_TaxID=6183;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Schmitz J., Symmons P., Dargatz H., Kunz W.;
 RL Submitted (FEB-1992) to the EMBL/GenBank/DBJ databases.
 DR EMBL: M81338; AAA29906.1;
 KW Myosin.
 FT NON_TER
 SQ SEQUENCE 148 AA; 17923 MW; C7EDA5A0BBE14DDA CRC64;

Query Match 53.8%; Score 49; DB 5; Length 148;
 Best Local Similarity 62.5%; Pred. No. 11;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
 QY 1 LKKRPMW 8
 :||| 1
 Db 30 VLKRPWM 37

RESULT 12
 ID 09Y806 PRELIMINARY; PRT; 298 AA.
 AC 09Y806:
 DT 01-NOV-1999 (TREMBLrel. 12, Created)
 DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
 DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
 DE HYPOTHEITICAL 33.7 KDA PROTEIN APE2577.
 GN APE2577.
 OS Aeropyrum pernix.
 CC Archaea; Crenarchaeota; Desulfurococcales; Desulfurococcales;
 CC Aeropyrum.
 NCBI_TaxID=56636;
 RN [1]
 RP SEQUENCE FROM N.A.

RC STRAIN-K1
 RX MEDLINE=99310339; PubMed=10382966;
 RA Kawarabayashi Y., Hino Y., Horikawa H., Yamazaki S., Halkawa Y.,
 RA Jin-no K., Takahashi M., Sekine M., Baba S.-I., Aikai A., Kosugi H.,
 RA Hosoyama A., Fukui S., Nagai Y., Nishijima K., Nakazawa H.,
 RA Takamiya M., Masuda S., Funahashi T., Tanaka T., Kudoh Y.,
 RA Yamazaki J., Kushiida N., Oguchi A., Aoki K.-I., Kubota K.,
 RA Nakamura Y., Nomura N., Sako Y., Kikuchi H.;
 RT "Complete genome sequence of an aerobic hyper-thermophilic
 RT crenarchaeon, Aeropyrum pernix K1."
 RL DNA Res. 6:83-101(1999).
 DR EMBL: AP000064; BAA81594.1;
 DR InterPro: IPR002787; DUF85.
 DR Pfam: PF01932; DUF85; 1.
 KW Hypothetical protein; Complete proteome.
 SQ SEQUENCE 298 AA; 33666 MW; FCB9C6EC93FE231 CRC64;

Query Match 53.8%; Score 49; DB 1; Length 298;
 Best Local Similarity 60.0%; Pred. No. 21;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
 QY 2 LKKRPMW 11
 :||| 1
 Db 102 LKRPWMWR 111

RESULT 13
 ID 019573 PRELIMINARY; PRT; 467 AA.
 AC 019573:
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-JUN-2000 (TREMBLrel. 14, Last annotation update)
 DE SIMILARITY TO 9 AMINO ACID REPEATS IN GALACTOSE SPECIFIC LECTINS.
 GN F1865.2.
 OS Caenorhabditis elegans.
 CC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
 CC Rhabditidae; Felodertinae; Caenorhabditis.
 NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=94150718; PubMed=7906398;
 RA Wilson R., Ainscough R., Anderson K., Baynes C., Berks M.,
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
 RA Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,
 RA Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,
 RA Jones M., Kershaw J., Kirsten J., Laister N., Latelle P.,
 RA Lightning J., Lloyd C., McMurray A., Mortimore B., O'Callaghan M.,
 RA Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Showkeen R.,
 RA Smaldon N., Smith A., Sonhammer E., Staden R., Sulston J.,
 RA Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K., Waterston R.,
 RA Watson A., Weinstock L., Wilkinson-Sproat J., Wohlman P.,
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans."
 RL Nature 368:32-38(1994).
 [2]

RP SEQUENCE FROM N.A.
RA Favellio T.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RA Waterston R.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL: U39855; AAA81082.1; -
KW lectin.
SQ SEQUENCE 467 AA; 63169 MW; 7D9BBAB61830431B CRC64;

Query Match 53.8%; Score 49; DB 5; Length 467;
Best Local Similarity 83.3%; Pred. No. 31;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 WPMWPMW 10
Db 201 WPMWPMW 206

RESULT 14
O26589 PRELIMINARY; PRT; 528 AA.
AC Q26589;
DT 01-NOV-1996 (TREMblrel. 01, Created)
DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE MYOSIN II HEAVY CHAIN (FRAGMENT).
OS Schistosoma mansoni (Blood fluke).
OC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
OC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
OC Schistosoma.
OX NCBI_TaxID=6183;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PUERTO RICAN;
RA Amory L.M.;
RL Submitted (APR-1992) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=PUERTO RICAN;
RX MEDLINE=93056536; Pubmed=1431131;
RA Soisson L.M., Masterston C.P., Tom T.D., McNally M.T., Lowell G.H.,
RA Strand M.;
RT "Induction of protective immunity in mice using a 62-kDa recombinant
fragment of a Schistosoma mansoni surface antigen.";
RL J. Immunol. 149:3612-3620(1992).
DR EMBL: X65591; CAA46548.1; -
DR HSSP: P08799; 1MMD.
DR InterPro: IPR001637; GlnA_adenyltn.
DR InterPro: IPR000048; IQ.
DR InterPro: IPR001609; myosin_head.
DR InterPro: IPR002928; Myosin_tail.
DR InterPro: IPR000533; Tropomyosin.
DR Pfam: PF00612; IQ; 1.
DR Pfam: PF00663; myosin_head; 1.
DR PRINTS: PR01576; Myosin_tail; 1.
DR PRINTS: PR00194; TROPOMYOSIN.
DR PRODOM: PD000355; myosin_head; 1.
DR SMART: SM00015; IQ; 1.
DR PROSITE: PS00182; GlnA_ADENYLATION; UNKNOWN_1.
DR PROSITE: PS50096; IQ; 1.
KW myosin.
FT NON_TER 1
SQ SEQUENCE 528 AA; 61622 MW; AF075D13EB249B4C CRC64;

Query Match 53.8%; Score 49; DB 5; Length 528;
Best Local Similarity 62.5%; Pred. No. 35;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
OY 1 ILKRWPMW 8

Db 106 VLRRWPMW 113

RESULT 15
O9TY57 PRELIMINARY; PRT; 528 AA.
AC O9TY57;
DT 01-MAY-2000 (TREMblrel. 13, Created)
DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE MYOSIN HEAVY CHAIN (FRAGMENT).
OS Schistosoma japonicum (Blood fluke).
OC Eukaryota; Metazoa; Platyhelminthes; Rhabditophora; Neodermata;
OC Trematoda; Digenea; Strigeidida; Schistosomatidae; Schistosomatidae;
OC Schistosoma.
OX NCBI_TaxID=6182;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CHINESE MAINLAND STRAIN;
RX MEDLINE=99144454; Pubmed=9990643;
RA Zhang Y.B., Taylor M.G., Bickle Q.D.;
RT "Schistosoma japonicum myosin: cloning, expression and vaccination
studies with the homologue of the S.mansoni myosin fragment ITV-5.";
RL Parasite Immunol. 20:583-594(1998).
DR EMBL: U55313; AAC82221.1; -
DR HSSP: P08799; 1MMD.
DR InterPro: IPR000048; IQ.
DR InterPro: IPR001609; myosin_head.
DR InterPro: IPR002928; Myosin_tail.
DR InterPro: IPR000533; Tropomyosin.
DR Pfam: PF00612; IQ; 1.
DR Pfam: PF00663; myosin_head; 1.
DR PRINTS: PR01576; Myosin_tail; 1.
DR PRINTS: PR00194; TROPOMYOSIN.
DR PRODOM: PD000355; myosin_head; 1.
DR SMART: SM00015; IQ; 1.
FT NON_TER 1
SQ SEQUENCE 528 AA; 61406 MW; C54A31F540F5EE05 CRC64;

Query Match 53.8%; Score 49; DB 5; Length 528;
Best Local Similarity 62.5%; Pred. No. 35;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 ILKRWPMW 8
Db 106 VLRRWPMW 113

Search completed: January 4, 2002, 08:47:24
Job time: 412 sec

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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:47 ; Search time 18.1 Seconds
(without alignments)
26.334 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKKMPWMPWRRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 100059 seqs, 36664827 residues
Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 10%
Listing first 45 summaries

Database : SwissProt_39.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the total score distribution, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	73	80.2	144	1 INDC_BOVIN	P33046 bos taurus
2	54	59.3	1173	1 VGL2_CVH22	P15423 human coron
3	49	53.8	492	1 ADRO_BOVIN	P08165 bos taurus
4	49	53.8	715	1 YD55_MYCTU	Q11025 mycobacteri
5	47	51.6	253	1 Y945_MYCTU	P71564 mycobacteri
6	46.5	51.1	1154	1 VGL2_IBVD2	P12222 avian infec
7	46.5	51.1	1162	1 VGL2_IBVB	P11223 avian infec
8	46.5	51.1	1162	1 VGL2_IBVK	P12650 avian infec
9	46.5	51.1	1162	1 VGL2_IBVM	P12651 avian infec
10	46.5	51.1	1163	1 VGL2_IBV6	P05135 avian infec
11	46	50.5	196	1 YA05_SCHPO	Q09677 schizosacch
12	45	49.5	397	1 MMU6_MYCTU	Q10773 mycobacteri
13	45	49.5	505	1 TREP_PSESS	P21689 pseudomonas
14	45	49.5	512	1 FEN2_YEAST	P25621 saccharomyc
15	45	49.5	964	1 MMU5_MYCTU	O53784 mycobacteri
16	45	49.5	967	1 MMU4_MYCTU	O53735 mycobacteri
17	45	49.5	968	1 MMU2_MYCTU	Q11171 mycobacteri
18	45	49.5	1108	1 CN3B_RAT	O63085 rattus norv
19	45	49.5	1225	1 VGL2_CVPR8	P27655 porcine res
20	45	49.5	1225	1 VGL2_CVPRM	P24413 porcine res
21	45	49.5	1235	1 VGL2_CVPMH	P11225 murine coro
22	45	49.5	1334	1 VGL2_CVMA5	P11224 murine coro
23	45	49.5	1353	1 VGL2_CVHOC	P36334 human coron
24	45	49.5	1363	1 VGL2_CVMB	P25190 bovine coro
25	45	49.5	1363	1 VGL2_CVBL5	P25191 bovine coro
26	45	49.5	1363	1 VGL2_CVBL5	P25192 bovine coro
27	45	49.5	1363	1 VGL2_CVBM	P15777 bovine coro
28	45	49.5	1363	1 VGL2_CVBO	P25193 bovine coro
29	45	49.5	1363	1 VGL2_CVBV	P25194 bovine coro
30	45	49.5	1376	1 VGL2_CVMA	P22432 murine coro
31	45	49.5	1376	1 VGL2_CVMC	Q02385 murine coro
32	45	49.5	1447	1 VGL2_CVPR	Q02167 porcine tira
33	45	49.5	1447	1 VGL2_CVPR	P07946 porcine tira

34	45	49.5	1447	1 VGL2_CVPR	Q01977 porcine tira
35	45	49.5	1449	1 VGL2_CVPR	P18450 porcine tira
36	45	49.5	1449	1 VGL2_CVPR	P33470 porcine tira
37	45	49.5	1451	1 VGL2_CVCAI	P36300 canine ente
38	45	49.5	1452	1 VGL2_CVCAI	P10033 feline inte
39	45	49.5	1452	1 VGL2_CVCAI	P08799 dictyostell
40	44	48.4	53	1 YDH3_PLAFS	P14589 plasmodium
41	44	48.4	151	1 YBHJ_ECOLI	P75709 escherichia
42	44	48.4	361	1 FURJ_HUMAN	P21217 homo sapien
43	44	48.4	372	1 FURJ_HUMAN	O19058 pan troglod
44	44	48.4	451	1 MENB_ECOLI	P37353 escherichia
45	44	48.4	535	1 YDWB_SCHPO	O13912 schizosacch

ALIGNMENTS

RESULT	1	STANDARD:	PRT:	144	AA.
ID	INDC_BOVIN				
AC	P33046:				
DT	01-OCT-1993 (Rel. 27, Created)				
DT	01-OCT-1993 (Rel. 27, Last sequence update)				
DT	01-NOV-1997 (Rel. 35, Last annotation update)				
DE	INDOLICIDIN PRECURSOR.				
OS	Bos taurus (Bovine).				
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;				
OX	Bovidae; Bovinae; Bos.				
OX	NCBI_TaxID=9913;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	TISSUE=Bone marrow;				
RX	MEDLINE=92392368; PubMed=1520337;				
RA	del Sal G., Storici P., Schneider C., Romeo D., Zanetti M.;				
RL	"CDNA cloning of the neutrophil bactericidal peptide indolicidin.";				
RL	Biochem. Biophys. Res. Commun. 187:467-472(1992).				
RN	[2]				
RP	SEQUENCE OF 131-143.				
RC	TISSUE=Neutrophils;				
RX	MEDLINE=92165771; PubMed=1537821;				
RA	Selsted M.E., Novotny M.J., Morris W.L., Tang Y.-Q., Smith W.;				
RT	Cullor J.S.;				
RT	"Indolicidin, a novel bactericidal tridecapeptide amide from				
RT	neutrophils.";				
RL	J. Biol. Chem. 267:4292-4295(1992).				
CC	- FUNCTION: POTENT MICROBICIDAL ACTIVITY, ACTIVE AGAINST				
CC	STAPHYLOCOCCUS AUREUS AND ESCHERICHIA COLI.				
CC	- TISSUE SPECIFICITY: LARGE GRANULES OF NEUTROPHILS.				
CC	- PTM: ELASTASE MIGHT BE RESPONSIBLE FOR ITS MATURATION.				
CC	- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.				
CC	-----				
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CC	or send an email to license@isb-sid.ch).				
CC	-----				
CC	EMBL: X67340; CAA47755.1; -				
DR	PIR: JCI222; JCI222.				
DR	PIR: A42387; A42387.				
DR	InterPro: IPR001894; Cathelicidin.				
DR	Pfam: PF00666; Cathelicidins; 1.				
DR	ProDom: PD001838; Cathelicidins; 1.				
DR	PROSITE: PS00946; CATHELICIDINS_1; 1.				
DR	PROSITE: PS00947; CATHELICIDINS_2; 1.				
KW	Antibiotic; Amidation; Signal.				
FT	SIGNAL	1	29		POTENTIAL.
FT	PROPEP	30	130		
FT	PEPTIDE	131	143		INDOLICIDIN.
FT	MOD_RES	30	30		PYRROLIDONE CARBOXYLIC ACID (BY

FT DISULFID 85 96 SIMILARITY).
FT CARBOHYD 107 124 BY SIMILARITY.
FT DISULFID 107 124 BY SIMILARITY.
FT MOD_RES 143 143 AMIDATION (G-144 PROVIDE AMIDE GROUP).
SQ SEQUENCE 144 AA; 16479 MW; E3B1CB5E5C09911 CRC64;

Query Match 80.2%; Score 73; DB 1; Length 144;
Best local Similarity 100.0%; Pred. No. 0.00094;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 KWPMPWR 12
DB 135 KWPMPWR 143

RESULT 2
VGL2_CVH22 STANDARD; PRT: 1173 AA.
ID VGL2_CVH22
AC P15423;

DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN).
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11137;

RP SEQUENCE FROM N.A.
RX MEDLINE=90264837; PubMed=2245367;
RA Raabe T., Schelle-Prinz B., Siddell S.G.;
RT "Nucleotide sequence of the gene encoding the spike glycoprotein of
human coronavirus HCoV 229E."
RT J. Gen. Virol. 71:1065-1073(1990).
CC -1- FUNCTION: THE PEPLOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION
AND IN SYNCYTIUM FORMATION.
CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.

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or send an email to license@isb-sib.ch).

CC EMBL: X16816; CAA34723.1; -.
DR PIR: A34766; VGIHHC.
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
KW Glycoprotein; Envelope protein; Transmembrane; Signal.
FT SIGNAL 1 15
FT CHAIN 16 1173 E2 GLYCOPROTEIN.
FT DOMAIN 16 1115 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 1116 1135 POTENTIAL.
FT DOMAIN 1136 1173 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 1136 1173
FT CARBOHYD 23 23 CYS-RICH.
FT CARBOHYD 23 23
FT CARBOHYD 62 98 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 98 98 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 147 147 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 171 171 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 176 176 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 220 220 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 243 243 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 326 326 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 333 333 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 440 440 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 464 464 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 518 518 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 538 538 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 542 542 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 568 568 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 581 581 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 587 587 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 663 663 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 671 671 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 930 930 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1015 1015 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1020 1020 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1037 1037 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1049 1049 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1061 1061 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1066 1066 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1076 1076 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1082 1082 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1096 1096 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 1173 AA; 128639 MW; B9CA9A1A796B3BD CRC64;

Query Match 59.3%; Score 54; DB 1; Length 1173;
Best local Similarity 85.7%; Pred. No. 2.5;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 KWPMPW 10
DB 1113 KWPMPW 1119

RESULT 3
ID ADRO_BOVIN STANDARD; PRT: 492 AA.
AC P08165;

DT 01-AUG-1988 (Rel. 08, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE NADPH:ADRENODOXIN OXIDOREDUCTASE, MITOCHONDRIAL PRECURSOR
(EC 1.18.1.2) (ADRENODOXIN REDUCTASE) (AR) (PERREROXIN-NADP(+)
REDUCTASE).

GN PDXR OR ADXR.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;

RP SEQUENCE FROM N.A., AND ALTERNATIVE SPLICING.

RX MEDLINE=94177140; PubMed=8130767;
RA Takata Y., Sagara Y., Kono A., Sekimizu K., Horiuchi T.;
RT "Gene structure of bovine adrenodoxin reductase.";
RL Biol. Pharm. Bull. 16:1200-1206(1993).

RN [2]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.

RX MEDLINE=88198050; PubMed=3448086;
RA Sagara Y., Takata Y., Miyata T., Hara T., Horiuchi T.;
RT "Cloning and sequence analysis of adrenodoxin reductase cDNA from
bovine adrenal cortex.";
RL J. Biochem. 102:1335-1336(1987).

RN [3]
RP SEQUENCE FROM N.A.

RX MEDLINE=87270696; PubMed=3038094;
RA Nonaka T., Murakami H., Yabusaki Y., Kuramitsu S., Kagamiyama H.,
Yamano T., Okamoto M.;
RT "Molecular cloning and sequence analysis of full-length cDNA for mRNA
of adrenodoxin oxidoreductase from bovine adrenal cortex.";
RL Biochem. Biophys. Res. Commun. 145:1239-1247(1987).

RP SEQUENCE FROM N.A.
RC TISSUE-Adrenal cortex;
RX MEDLINE=89170752; PubMed=2924777;
RA Hanukoglu I., Gutfinger T.;

RT "cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites in oxidoreductases.";
 RL Eur. J. Biochem. 180:479-484(1989).
 RN [5]
 RP SEQUENCE OF N-TERMINUS, AND PARTIAL SEQUENCE.
 RX TISSUE-Adrenal cortex;
 RX MEDLINE=88082777; PubMed=3691502;
 RA Hanukoglu I., Gutfinger T., Hanin M., Shively J.E.;
 RT "Isolation of a cDNA for adrenodoxin reductase (ferredoxin-NADP+ reductase). Implications for mitochondrial cytochrome P-450 systems.";
 RL Eur. J. Biochem. 169:449-455(1987).
 RN [6]
 RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS) OF 33-492.
 RX TISSUE-Adrenal gland;
 RX MEDLINE=99299392; PubMed=10369776;
 RA Ziegler G.A., Vornrhein C., Hanukoglu I., Schulz G.E.;
 RT "The structure of adrenodoxin reductase of mitochondrial P450 systems: election transfer for steroid biosynthesis.";
 RL J. Mol. Biol. 289:981-990(1999).
 CC -1- FUNCTION: SERVES AS THE FIRST ELECTRON TRANSFER PROTEIN IN ALL THE MITOCHONDRIAL P450 SYSTEMS, INCLUDING CHOLESTEROL SIDE CHAIN CLEAVAGE IN ALL STEROIDGENIC TISSUES, STEROID 11-BETA HYDROXYLATION IN THE ADRENAL CORTEX, 25-OH-VITAMIN D3-24 HYDROXYLATION IN THE KIDNEY, AND STEROL C-27 HYDROXYLATION IN THE LIVER.
 CC -1- CATALYTIC ACTIVITY: REDUCED ADRENODOXIN + NADP(+) = OXIDIZED ADRENODOXIN + NADPH.
 CC -1- COFACTOR: FAD.
 CC -1- PATHWAY: CHOLESTEROL SIDE-CHAIN-CLEAVAGE SYSTEM.
 CC -1- SUBUNIT: MONOMER.
 CC -1- SUBCELLULAR LOCATION: MITOCHONDRIAL MATRIX.
 CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS; A SHORT FORM (SHOWN HERE) AND A LONG FORM; ARE PRODUCED BY ALTERNATIVE SPLICING. THE LONG FORM REPRESENTS 10-20% OF ALL ADRENODOXIN REDUCTASE MRNA. AND SEEMS TO BE INACTIVE.
 CC -----
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 CC -----
 DR EMBL: M17029; AAA0362.1; -;
 DR EMBL: D00211; BA00150.1; -;
 DR EMBL: X13736; CA032002.1; -;
 DR PIR: A29604; A29604.
 DR PIR: J50390; J50390.
 DR PIR: S03558; S03558.
 DR PIR: J70751; J70751.
 DR PDB: 1CJC; 12-APR-99.
 DR PDB: 1E1L; 02-JUN-00.
 DR InterPro: IPR000759; Adrnkx_reductase.
 DR PRINTS: PR00419; ADXRDVASE.
 KW Electron transport; Oxidoreductase; Flavoprotein; NADP; FAD;
 KW Mitochondrion; Transit peptide; Alternative splicing; 3D-structure.
 FT TRANSIT 1 32
 FT CHAIN 33 492
 FT VARSPLIC 204 204
 FT CONFLICT 77 77
 FT CONFLICT 81 94
 FT CONFLICT 124 128
 FT CONFLICT 268 268
 FT CONFLICT 317 318
 FT CONFLICT 333 333
 FT CONFLICT 341 352
 FT SEQUENCE 492 AA; 54338 MW; E68F6F5D18F53131 CRC64;
 Query Match 53.8%; Score 49; DB 1; Length 492;
 Best Local Similarity 43.3%; Pred. No. 5.4;

Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 5 WPMPWP 10
 Db 6 WRMPWP 11
 RESULT 4
 ID YD55_MYCTU STANDARD; PRT; 715 AA.
 AC Q11025;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DE 20-AUG-2001 (Rel. 40, Last annotation update)
 DE HYPOTHETICAL 78.2 KDA PROTEIN RV1355C.
 GN RV1355C OR MT1398 OR MYCY02B10.19C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D., Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F., Badcock K., Basham D., Brown D., Chillingworth T., Connor R., Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S., Hornsby T., Jagels K., Krogh A., McLean A., Moule S., Murphy L., Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J., Rutter S., Seeger K., Skelton S., Squares S., Squares R., Sultun J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.";
 RT Nature 393:537-544(1998).
 RL [2]
 RN SEQUENCE FROM N.A.
 RP STRAIN=CDC 1551 / Oshkosh;
 RC Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O., Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E., Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L., Ralcher A., Ustebach T., Weidman J., Khouri H., Gill J., Mikula A., Bishtal W.;
 RA "Whole genome comparison of Mycobacterium tuberculosis clinical and laboratory strains.";
 RT Submitted (Apr-2001) to the EMBL/Genbank/DBSJ databases.
 RL [3]
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 CC -----
 DR EMBL: Z75555; CA099988.1; -;
 DR EMBL: AE007012; AAK45661.1; ALT_INIT.
 DR TIGR: MT1398; -;
 DR TubercuList: RV1355C; -;
 DR InterPro: IPR000594; Thif_family.
 DR Pfam: PF00899; Thif_family; 1.
 KW Hypothetical protein; Complete proteome.
 FT SEQUENCE 715 AA; 78181 MW; 455495248A56041C CRC64;
 Query Match 53.8%; Score 49; DB 1; Length 715;
 Best Local Similarity 60.0%; Pred. No. 7.6;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
 QY 3 KKMPMPMP 12
 Db 64 KRWMPMP 73

Y	6	PWMPW	10	51.6%	Score 47;	DB 1;	Length 253;
db	230	PWMPW	234	Best Local Similarity 100.0%;	Pred. No. 5.4;		
	Matches	5;	Conservative	0;	Mismatches	0;	Indels
						0;	Gaps
0Y	6	PWMPW	10	51.6%	Score 47;	DB 1;	Length 253;
	230	PWMPW	234	Best Local Similarity 100.0%;	Pred. No. 5.4;		
	Matches	5;	Conservative	0;	Mismatches	0;	Indels
						0;	Gaps

ID	VL2	IBVD2	STANDARD:	PRT:	1154 AA.
AC	P12722:	Q66176; Q66177;			
DT	01-OCT-1989	(Rel. 12, Created)			
DT	01-OCT-1989	(Rel. 12, Last sequence update)			
DT	20-AUG-2001	(Rel. 40, Last annotation update)			
DE	E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN)	(PEPLOMER PROTEIN)			
DE	[CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].				
GN	S.				
OS	Avian infectious bronchitis virus (strain D274) (IBV).				
OC	viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;				
OC	Coronaviridae; Coronaviruses.				
NCBI	taxid=11124;				
RM	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=89386000; PubMed=2550899;				
RA	Jordi B.J.A.M., Kreners D.A.W.M., Kusters H.G., van der Zeijst B.A.M.;				
RT	"Nucleotide sequence of the gene coding for the peplomer protein (= spike protein) of infectious bronchitis virus, strain D274."				
RL	Nucleic Acids Res. 17:6726-6726(1989).				
CC	-I- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.				
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CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -				
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CC	use by non-profit institutions as long as its content is in no way				
CC	modified and this statement is not removed. Usage by and for commercial				
CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/				
CC	or send an email to license@sib-sib.ch).				
CC	-----				
DR	EMBL, X15832: CAA33837.1; -				
DR	PIR, A34300; VGHIIB.				
DR	InterPro: IPR002551; Corona_S1.				
DR	InterPro: IPR002552; Corona_S2.				
DR	Pfam: PF01600; Corona_S1; 1.				
DR	Pfam: PF01601; Corona_S2; 1.				
KW	Glycoprotein; Envelope protein; Transmembrane; Signal.				
FT	SIGNAL	1			
FT	CHAIN	19	1154		
FT	CHAIN	19	538		
FT	CHAIN	539	1154		
FT	DOMAIN	1121	1138		
FT	CARBOHYD	23	23		
FT	CARBOHYD	74	74		
FT	CARBOHYD	102	102		
FT	CARBOHYD	139	139		
FT	CARBOHYD	145	145		
FT	CARBOHYD	164	164		
FT	CARBOHYD	179	179		
FT	CARBOHYD	213	213		
FT	CARBOHYD	238	238		
FT	CARBOHYD	248	248		
FT	CARBOHYD	265	265		
FT	CARBOHYD	272	272		
FT	CARBOHYD	277	277		
FT	CARBOHYD	307	307		
FT	CARBOHYD	426	426		
FT	CARBOHYD	448	448		
FT	CARBOHYD	514	514		
FT	CARBOHYD	531	531		
FT	CARBOHYD	543	543		
FT	CARBOHYD	580	580		
FT	CARBOHYD	592	592		
FT	CARBOHYD	670	670		
FT	CARBOHYD	677	677		
FT	CARBOHYD	948	948		
FT	CARBOHYD	961	961		
FT	CARBOHYD	980	980		
FT	CARBOHYD	1015	1015		
FT	CARBOHYD	1039	1039		
FT	CARBOHYD	1052	1052		
FT	CARBOHYD	1075	1075		

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SQ SEQUENCE 1154 AA; 127502 MW; D79F37AF89F1A37F CRC64;
Query Match
Best Local Similarity 51.1%; Score 46.5; DB 1; Length 1154;
Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK---KMPMPW 10
   ||| |||||
Db 1086 ILKTYIKMPWYVW 1098

RESULT 7
VGL2_IBVK STANDARD; PRT; 1162 AA.
ID AC P12650;
DT 01-JUL-1989 (Rel. 11, Created)
DT 01-JUL-1989 (Rel. 11, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)
DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].
GN S.
OS Avian infectious bronchitis virus (strain Beaudette) (IBV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11122;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=85159540; PubMed=2984314;
RA Binns M.M., Boursnell M.E.G., Cavanagh D., Pappind D.J.C.,
RA Brown T.D.K.;
RT "Cloning and sequencing of the gene encoding the spike protein of the
RT coronavirus IBV";
RL J. Gen. Virol. 66:719-726(1985).
[2]
RN SEQUENCE FROM N.A.
RX MEDLINE=87085499; PubMed=3025348;
RA Binns M.M., Boursnell M.E.G., Tomley F.M., Brown D.K.;
RT "Comparison of the spike precursor sequences of coronavirus IBV
RT strains M1 and 6/82 with that of IBV Beaudette.";
RL J. Gen. Virol. 67:2825-2831(1986).
CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
CC -----
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CC or send an email to license@sib-sib.ch).
CC -----
CC EMBL; M95169; AAA70235.1; -
DR EMBL; X02342; CAA26201.1; -
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
KW Glycoprotein; Envelope protein; Transmembrane; Signal.
FT SIGNAL 1
FT CHAIN 19 1162 E2 GLYCOPROTEIN.
FT CHAIN 19 537 SPIKE PROTEIN S1.
FT CHAIN 1120 1137 SPIKE PROTEIN S2.
FT DOMAIN 51 1137 CYS-RICH.
FT CARBOHYD 77 79 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).

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FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1038 1038 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 1162 AA; 128046 MW; 0BAAD58113C8EBD5 CRC64;

Query Match
Best Local Similarity 51.1%; Score 46.5; DB 1; Length 1162;
Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILK---KMPMPW 10
   ||| |||||
Db 1085 ILKTYIKMPWYVW 1097

RESULT 8
VGL2_IBVK STANDARD; PRT; 1162 AA.
ID AC P12650;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)
DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].
GN S.
OS Avian infectious bronchitis virus (strain KB8523) (IBV).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=11126;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88306251; PubMed=2841803;
RA Sutoh S., Sato S., Okabe T., Nakai M., Sasaki N.;
RT "Cloning and sequencing of genes encoding structural proteins of
RT avian infectious bronchitis virus.";
RL Virology 165:589-595(1988).
CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS
CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
CC -----
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CC or send an email to license@sib-sib.ch).
CC -----
CC EMBL; M21515; AAA66578.1; -
DR PIR; B93249; VG1HAK.
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
KW Glycoprotein; Envelope protein; Transmembrane; Signal.
FT SIGNAL 1
FT CHAIN 19 1162 E2 GLYCOPROTEIN.
FT CHAIN 19 537 SPIKE PROTEIN S1.

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FT CHAIN 538 1162 SPIKE PROTEIN S2.
FT DOMAIN 1120 1137 CYS-RICH.
FT CARBOHYD 51 81. N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 77 77. N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 271 271 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 676 676 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1058 1058 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 1162 AA; 128537 MW; 2299036B3597EAB8F CRC64;

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Query Match .51.1%; Score 46.5; DB 1; Length 1162;

Best Local Similarity 61.5%; Pred. No. 26; Mismatches 1; Indels 3; Gaps 1;

Matches 8; Conservative 1; Indels 3; Gaps 1;

Qy 1 ILK---KMPMPW 10
 ||| ||||: |
 Db 1085 ILKTYIKMPWYV 1097

RESULT 9

VG12_IBVM STANDARD; PRT; 1162 AA.

AC P12651; 01-OCT-1989 (Rel. 12, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)

DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].

GN S.

OS Avian infectious bronchitis virus (strain M41) (IBV).

OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.

OX NCBI_TaxID=11127;

RN [1]

RP MEDLINE=87021475; PubMed=2429473;

RX Niesters H.G.M., Lenstra J.A., Spaan W.J.M., Zijderfeld A.J.,

RA Bleumink-Pluym N.M.C., Hong F., Van Scharrenburg G.J.M.,

RA Horzinek M.C., van der Zeijst B.A.M.;

RT "The peplomer protein sequence of the M41 strain of coronavirus IBV

and its comparison with Beaudette strains.";

RL Virus Res. 5:253-263(1986).

CC -1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS

CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.

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CC -----

DR EMBL; M21883; AAA65575.1; -;

DR EMBL; A24863; CA01736.1; -;

DR PIR; S07421; S07421.

DR InterPro; IPR002551; Corona_S1.

DR InterPro; IPR002552; Corona_S2.

DR Pfam; PF01601; Corona_S1; 1.

DR Pfam; PF01601; Corona_S2; 1.

KW Glycoprotein; Envelope protein; Transmembrane; Signal.

FT SIGNAL 1 18

FT CHAIN 19 1162 E2 GLYCOPROTEIN.

FT CHAIN 19 537 SPIKE PROTEIN S1.

FT CHAIN 538 1162 SPIKE PROTEIN S2.

FT DOMAIN 1120 1137 CYS-RICH.

FT CARBOHYD 51 51 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 77 77 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 103 103 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 144 144 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 163 163 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 178 178 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 212 212 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 237 237 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 264 264 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 271 271 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 276 276 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 306 306 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 425 425 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 447 447 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 513 513 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 530 530 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 579 579 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 591 591 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 669 669 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 714 714 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 947 947 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 960 960 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 979 979 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1014 1014 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1038 1038 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1051 1051 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 1074 1074 N-LINKED (GLCNAC. . .) (POTENTIAL).

SQ SEQUENCE 1162 AA; 128077 MW; 3C9CC70938492DDA CRC64;

Query Match .51.1%; Score 46.5; DB 1; Length 1162;

Best Local Similarity 61.5%; Pred. No. 26; Mismatches 1; Indels 3; Gaps 1;

Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

Qy 1 ILK---KMPMPW 10
 ||| ||||: |
 Db 1085 ILKTYIKMPWYV 1097

RESULT 10

VG12_IBV6 STANDARD; PRT; 1163 AA.

AC P05135; 13-AUG-1987 (Rel. 05, Created)

DT 13-AUG-1987 (Rel. 05, Last sequence update)

DE E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN)

DE [CONTAINS: SPIKE PROTEIN S1; SPIKE PROTEIN S2].

GN S.

OS Avian infectious bronchitis virus (strain 6/82) (IBV).

OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.

OX NCBI_TaxID=11121;

RN [1]

RP SEQUENCE FROM N.A.
 RA MEDLINE-87085499; PubMed-3025348;
 RT Bins M.M., Bournell M.E.G., Tomley F.M., Brown D.K.;
 RT "Comparison of the spike precursor sequences of coronavirus IBV
 RT strains M41 and 6/82 with that of IBV Beaudette.";
 RL J. Gen. Virol. 67:2825-2831(1986).
 CC -1- FUNCTION: THE PEPLIMER PROTEIN MEDIATES THE BINDING OF VIRIONS
 CC TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION.
 CC -----
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 CC -----
 CC EMBL: X04723; CAA28432.1; -
 DR InterPro: IPR002551; Corona_S1.
 DR InterPro: IPR002552; Corona_S2.
 DR Pfam: PF01600; Corona_S1; 1.
 DR Pfam: PF01601; Corona_S2; 1.
 KW Glycoprotein; Transmembrane; Signal.
 FT SIGNAL 1 18
 FT CHAIN 19 1163 E2 GLYCOPROTEIN.
 FT CHAIN 19 538 SPIKE PROTEIN S1.
 FT CHAIN 539 1163 SPIKE PROTEIN S2.
 FT DOMAIN 1121 1138 CYS-RICH.
 FT CARBOHYD 23 23 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 51 51 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 74 74 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 102 102 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 139 139 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 145 145 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 164 164 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 179 179 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 213 213 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 238 238 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 248 248 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 265 265 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 272 272 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 277 277 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 307 307 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 426 426 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 448 448 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 514 514 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 531 531 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 543 543 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 580 580 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 592 592 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 670 670 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 677 677 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 948 948 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 961 961 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 980 980 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1015 1015 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1039 1039 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1052 1052 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 1075 1075 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 1163 AA; 128684 MW; 8FE344CF2995A78C CRC64;

Query Match 51.1%; Score 46.5; DB 1; Length 1163;
 Best Local Similarity 61.5%; Pred. No. 26;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

OY 1 ILKWP-MW---PMRKP 10
 DB 1086 ILKTIKMPYVW 1098

RESULT 11
 YAO5_SCHPO

ID YAO5_SCHPO STANDARD; PRT; 196 AA.
 AC 009677;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 01-NOV-1995 (Rel. 32, Last annotation update)
 DE HYPOTHETICAL 22.1 KDA PROTEIN C5H10.05C IN CHROMOSOME I.
 GN SPAC5H10.05C.
 OS Schizosaccharomyces pombe (fission yeast).
 OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;
 OC Schizosaccharomycetes.
 OX NCBI_TaxID=4896;
 OX (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN-972;
 RA Connor R., Churche C.M., Barrell B.G., Rajandream M.A., Walsh S.V.;
 RL Submitted (MAY-1995) to the EMBL/Genbank/DBJ databases.
 CC -1- SIMILARITY: STRONG, TO BACTERIAL MODULATOR OF DRUG ACTIVITY B
 CC (MDAB).
 CC -----
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 CC -----
 CC EMBL: Z49811; CAA89955.1; -
 DR InterPro: IPR003680; NADHdh_2.
 DR Pfam: PF02525; NADHdh_2; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 196 AA; 22104 MW; 436764DA9E26074C CRC64;

Query Match 50.5%; Score 46; DB 1; Length 196;
 Best Local Similarity 47.1%; Pred. No. 5.9;
 Matches 8; Conservative 3; Mismatches 2; Indels 4; Gaps 2;

OY 1 ILKWP-MW---PMRKP 13
 DB 62 ILYWPGWMMGPWKLR 78

RESULT 12
 MH16_MYCTU STANDARD; PRT; 397 AA.
 AC Q10773;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MHP16.
 GN MHP16 OR RV1557 OR MHP1608 OR MTCY48.08C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1773;
 OX (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN-H37RV;
 RX MEDLINE-98295987; PubMed-9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekle A.F.,
 RA Badoock K., Basham D., Brown D., Chillingworth T., Connor R.,
 RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holtroyd S.,
 RA Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
 RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 RA Rutter S., Seeger K., Skellern S., Squares S., Squares R.,
 RA Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the
 RA complete genome sequence.";
 RL Nature 393:537-544(1998).
 RN [2]

```

RP SEQUENCE FROM N.A.
RC STRAIN=CDC 1551 / Oshkosh;
RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
RA Peterson J., Deboy R., Dodson R., Gwin M.L., Haft D., Hickey E.,
RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
RA Delcher A., Uterback T., Weidman J., Kouri H., Gill J., Mikula A.,
RA Biswal W.;
RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
RT laboratory strains."
RL Submitted (Apr-2001) to the EMBL/Genbank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
CC -1- SIMILARITY: BELONGS TO THE MML FAMILY.
CC -----
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CC -----
DR EMBL; Z74020; CAA98334.1; -
DR EMBL; AE007027; AAK45875.1; -
DR TIGR; MT1608; -
DR TubercuList; Rv1557; -
KW Hypothetical protein; Transmembrane; Complete proteome.
FT TRANSMEM 161 181 POTENTIAL.
FT TRANSMEM 190 210 POTENTIAL.
FT TRANSMEM 214 234 POTENTIAL.
FT TRANSMEM 242 262 POTENTIAL.
FT TRANSMEM 293 313 POTENTIAL.
FT TRANSMEM 330 350 POTENTIAL.
SQ SEQUENCE 397 AA; 42421 MW; 678DC86E24472BF4 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 397;
Best Local Similarity 54.5%; Pred. No. 15;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 ILKWPWPWR 11
Db 348 LLGRWFWPQR 358

RESULT 13
TRPE_PSESS STANDARD; PRT; 505 AA.
AC P21689;
DT 01-MAY-1991 (Rel. 18, Created)
DT 01-MAY-1991 (Rel. 18, Last sequence update)
DT 30-MAY-2000 (Rel. 39, Last annotation update)
DE ANTHRANILATE SYNTHASE COMPONENT I (EC 4.1.3.27).
GN TRPE.
OS Pseudomonas syringae (pv. savastanoi).
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
OC Pseudomonas.
OX NCBI_TaxID=29438;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91100331; PubMed=1987141;
RA da Costa E., Silva O., Kosuge T.;
RT "Molecular characterization and expression analysis of the
RT anthranilate synthase gene of Pseudomonas syringae subsp.
RT savastanoi."
RL J. Bacteriol. 173:463-471(1991).
CC -1- CATALYTIC ACTIVITY: CHORISMATE + L-GLUTAMINE = ANTHRANILATE +
CC PYRUVATE + L-GLUTAMATE.
CC -1- PATHWAY: FIRST STEP IN BIOSYNTHESIS OF TRYPTOPHAN.
CC -1- SUBUNIT: TRIMER OF TWO COMPONENTS I AND TWO COMPONENTS II (BY
CC SIMILARITY).
CC -1- MISCELLANEOUS: COMPONENT I CATALYZES THE FORMATION OF ANTHRANILATE
CC USING AMMONIA RATHER THAN GLUTAMINE, WHEREAS COMPONENT II PROVIDES
CC GLUTAMINE AMIDOTRANSFERASE ACTIVITY.

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CC -1- SIMILARITY: BELONGS TO THE ANTHRANILATE SYNTHASE COMPONENT I
CC FAMILY.
CC -----
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CC -----
DR EMBL; M55911; AAA26016.1; -
DR PIR; A39128; A39128.
DR InterPro; IPR000350; Chorismate_bind.
DR Pfam; PF00425; chorismate_bind; 1.
DR PRINTS; PR00095; ANTSYNTHASE1.
DR Prodom; PD000779; Chorismate_bind; 1.
KW Tryptophan biosynthesis; Lyase.
SQ SEQUENCE 505 AA; 56084 MW; A38E81931331F6B8 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 505;
Best Local Similarity 71.4%; Pred. No. 19;
Matches 5; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 7 WWPWRK 13
Db 485 WWPWR 491

RESULT 14
FEN2_YEAST
ID FEN2_YEAST STANDARD; PRT; 512 AA.
AC P25621;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE PROBABLE TRANSPORTER FEN2.
GN FEN2 OR YCR028C OR YCR28C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RA Cederberg H., Hohmann S., Schaaff-gerstenschlaeger I., Huse K.,
RA Zimmermann F.K.; (1992) to the EMBL/Genbank/DBJ databases.
RL Submitted (MAR-1992) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=93070619; PubMed=1332309;
RA Carbone M.L.A., Panzeri L., Falconi M.M., Carcano C., Plevani P.,
RA Lucchini G.;
RT "Nucleotide sequence of 9.2 kb left of CRI1 on yeast chromosome III
RT from strain AB872: evidence for a ty insertion and functional
RT analysis of open reading frame YCR28."
RL Yeast 8:805-812(1992).
RN [3]
RP SIMILARITY TO DAL5 FAMILY.
RX MEDLINE=94147996; PubMed=8313894;
RA Koonin E.V., Bork P., Sander C.;
RT "Yeast chromosome III: new gene functions."
RL EMBO J. 13:493-503(1994).
RN [4]
RP CHARACTERIZATION.
RX MEDLINE=96367594; PubMed=8771708;
RA Marcireau C., Joets J., Poussel D., Guilloton M., Karst F.;
RT "FEN2: a gene implicated in the catabolite repression-mediated
RT regulation of ergosterol biosynthesis in yeast."
RL Yeast 12:531-539(1996).
CC -1- FUNCTION: INVOLVED IN THE CATABOLITE REPRESSION-MEDIATED
CC REGULATION OF ERGOSTEROL BIOSYNTHESIS AND IN FENPROPIOMORPH
CC RESISTANCE.

```

CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE ATLANTOATE PERMEASE FAMILY.
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 CC -----
 CC EMBL: X59720; CAA42320.1;
 CC DR PIR: S19439; S19439.
 CC DR PIR: S25336; S25336.
 CC DR SGD: S0000623; FEN2.
 CC KW Transmembrane; Transport.
 CC FT TRANSMEM 28 48
 CC FT TRANSMEM 80 100 POTENTIAL.
 CC FT TRANSMEM 103 123 POTENTIAL.
 CC FT TRANSMEM 133 153 POTENTIAL.
 CC FT TRANSMEM 165 185 POTENTIAL.
 CC FT TRANSMEM 199 219 POTENTIAL.
 CC FT TRANSMEM 272 292 POTENTIAL.
 CC FT TRANSMEM 313 333 POTENTIAL.
 CC FT TRANSMEM 343 363 POTENTIAL.
 CC FT TRANSMEM 373 393 POTENTIAL.
 CC FT TRANSMEM 402 422 POTENTIAL.
 CC FT TRANSMEM 435 455 POTENTIAL.
 CC FT CONFLICT 104 104 W -> V (IN REF. 2).
 CC SQ SEQUENCE 512 AA; 58256 MW; 361942E74C62B384 CRC64;

Query Match 49.5%; Score 45; DB 1; Length 512;
 Best Local Similarity 62.5%; Pred. No. 20;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILKKWMPW 8
 Db 268 VLKRWMMW 275
 RESULT 15
 MML5_MYCTU STANDARD; PRT; 964 AA.
 ID MML5_MYCTU STANDARD; PRT; 964 AA.
 AC 053784;
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML5.
 GN MML5 OR RV0676C OR MT0705 OR MT040.04C.
 OS Mycobacterium tuberculosis.
 CC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 CC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
 CC NCBI_TaxID=1773;
 OX 111
 RN SEQUENCE FROM N.A.
 RP STRAIN=H37RV;
 RC MEDLINE=98295987; PubMed=9634230;
 RX Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 RA Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F.,
 RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 RA Davies R., Devlin K., Feltwell T., Gentles S., Hamlin N., Holroyd S.,
 RA Hornsby T., Jagels K., Kiroch A., McLean J., Moule S., Murphy L.,
 RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 RA Ruter S., Seeger K., Skelton S., Squares S., Squares R.,
 RA Sulston J.E., Taylor K., Whitehead S., Barrett B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 RN 121
 RP SEQUENCE FROM N.A.
 RC STRAIN=CDC 1551 / Oshkosh;
 RX Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,

RA Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
 RA Kolonay J.F., Nelson W.C., Umayan L.A., Ermolaeva M.D., Salzberg S.L.,
 RA Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A.,
 RA Bishai W.;
 RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RT laboratory strains."
 RT Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML5 FAMILY.
 CC -----
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 CC -----
 CC EMBL: AL021943; CAA17459.1;
 CC DR EMBL: AE006964; AAK44930.1;
 CC DR TIGR: MT0705;
 CC DR TubercuList; RV0676C;
 CC DR InterPro; IPR001036; ACR_tran.
 CC DR PRINTS; PR00702; ACRIFLAVINRP.
 CC KW Hypothetical protein; Transmembrane; Complete proteome.
 CC FT TRANSMEM 31 51 POTENTIAL.
 CC FT TRANSMEM 203 223 POTENTIAL.
 CC FT TRANSMEM 230 250 POTENTIAL.
 CC FT TRANSMEM 255 275 POTENTIAL.
 CC FT TRANSMEM 302 322 POTENTIAL.
 CC FT TRANSMEM 340 360 POTENTIAL.
 CC FT TRANSMEM 389 409 POTENTIAL.
 CC FT TRANSMEM 745 765 POTENTIAL.
 CC FT TRANSMEM 774 794 POTENTIAL.
 CC FT TRANSMEM 803 823 POTENTIAL.
 CC FT TRANSMEM 826 846 POTENTIAL.
 CC FT TRANSMEM 880 900 POTENTIAL.
 CC FT TRANSMEM 901 921 POTENTIAL.
 CC FT CONFLICT 948 948 I -> V (IN REF. 2).
 CC SQ SEQUENCE 964 AA; 104784 MW; B7C945940A1176BD CRC64;

Query Match 49.5%; Score 45; DB 1; Length 964;
 Best Local Similarity 66.7%; Pred. No. 35;
 Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 ILKKWMPW 9
 Db 932 LLGKWFMP 940
 Search completed: January 4, 2002, 08:47:48
 Job time: 406 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:41:31 ; Search time 27.18 Seconds
(without alignments)
36,434 Million cell updates/sec

Title: US-09-444-281-35
Perfect score: 91
Sequence: 1 ILKKWMPWMPRRK 13

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues

Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: p1r1:
2: p1r2:
3: p1r3:
4: p1r4:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	73	80.2	144	JC1222	indolicidin precursor
2	54	59.3	1173	1_VG1HHC	E2 glycoprotein pr
3	51	56.0	289	2_T12505	hypothetical prote
4	49	53.8	298	2_B72492	hypothetical prote
5	49	53.8	498	1_JT0751	ferredoxin--NADP+
6	49	53.8	527	2_S33068	myosin heavy chain
7	49	53.8	715	2_B70741	probable myo prot
8	49	53.8	1940	2_A59287	myosin heavy chain
9	48	52.7	111	2_T29295	hypothetical prote
10	47.5	52.2	114	2_T36208	hypothetical prote
11	47	51.6	248	2_S23449	NADH oxidase (H2O2
12	47	51.6	253	2_G70715	histidinol-phospha
13	46.5	51.1	352	2_S77354	peplomeric polypro
14	46.5	51.1	621	2_S37664	peplomeric polypro
15	46.5	51.1	630	2_S37663	peplomeric polypro
16	46.5	51.1	1154	1_VG1HHC	E2 glycoprotein pr
17	46.5	51.1	1162	1_VG1HHC	E2 glycoprotein pr
18	46.5	51.1	1162	2_S07421	E2 glycoprotein pr
19	46.5	51.1	1162	2_S14939	E2 glycoprotein pr
20	46.5	51.1	1162	2_S14940	E2 glycoprotein pr
21	46	50.5	196	2_S55483	modulator of drug
22	46	50.5	617	2_T22175	hypothetical prote
23	46	50.5	623	2_T22177	hypothetical prote
24	46	50.5	1333	2_S65812	RNA-directed DNA p
25	45	49.5	273	2_F82646	monofunctional bio
26	45	49.5	276	2_E83161	probable short-cha
27	45	49.5	397	2_B70763	probable membrane
28	45	49.5	448	2_H72376	hypothetical prote
29	45	49.5	505	2_A39128	anthranilate synth

30	45	49.5	512	2_S19439	probable membrane
31	45	49.5	964	2_E70826	probable membrane
32	45	49.5	967	2_C70831	probable mmp14 pro
33	45	49.5	968	2_F70746	probable mmp12 pro
34	45	49.5	1108	2_A48508	cyclic-nucleotide
35	45	49.5	1225	2_S24284	E2 glycoprotein pr
36	45	49.5	1235	2_A36607	E2 glycoprotein pr
37	45	49.5	1235	2_VG1HHC	E2 glycoprotein pr
38	45	49.5	1324	1_VG1H59	E2 glycoprotein pr
39	45	49.5	1353	1_J02168	E2 glycoprotein pr
40	45	49.5	1361	2_S29998	surface protein
41	45	49.5	1362	2_A37474	surface glycoprote
42	45	49.5	1363	1_VG1HHC	E2 glycoprotein pr
43	45	49.5	1363	1_VG1H59	E2 glycoprotein pr
44	45	49.5	1363	1_VG1H59	E2 glycoprotein pr
45	45	49.5	1363	1_VG1H59	E2 glycoprotein pr

ALIGNMENTS

RESULT 1
JC1222
indolicidin precursor - bovine
N:Alternate names: antimicrobial peptide
C:Species: Bos primigenius taurus (cattle)
C:Date: 10-Sep-1999 #sequence,revision 10-Sep-1999 #text,change 10-Sep-1999
C:Accession: JC1222; A42387; S25664
R:del Sal, G.; Storici, P.; Schneider, C.; Romeo, D.; Zanetti, M.
Biochem. Biophys. Res. Commun. 187, 467-472, 1992
A:Title: cDNA cloning of the neutrophil bactericidal peptide indolicidin.
A:Reference number: JC1222; WUID:92392368
A:Accession: JC1222
A:Molecule type: mRNA
A:Residues: 1-144 <SAL>
A:Cross-references: EMBL:X67340; NID:9462; PIDN:CAA47755.1; PID:9463
A:Experimental source: bone marrow
J:Seasted, M.E.; Novotny, M.J.; Morris, W.L.; Tang, Y.Q.; Smith, W.; Cullor, J.S.
J. Biol. Chem. 267, 4292-4295, 1992
A:Title: Indolicidin, a novel bactericidal tridecapeptide amide from neutrophils.
A:Reference number: A42387; WUID:92165771
A:Accession: A42387
A:Molecule type: protein
A:Residues: 131-143 <SEL>
A:Experimental source: neutrophils
A:Note: sequence extracted from NCBI backbone (NCBI:83840)
C:Superfamily: cathelin; cystatin homology
C:Keywords: amidated carboxyl end
F:1-29/Domain: signal sequence #status predicted <SIG>
F:30-129/Domain: cystatin homology <CYS>
F:131-143/Product: propeptide #status predicted <PRO>
F:131-143/Product: indolicidin #status experimental <MNT>
F:143/Modified site: amidated carboxyl end (Arg) (amide in mature form from following

Query Match 80.2% Score 73; DB 1; Length 144;
Best Local Similarity 100.0% Pred. No. 0.0027;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 KWPMPWMPRR 12
Db 135 KWPMPWMPRR 143

RESULT 2
VG1HHC
E2 glycoprotein precursor - human coronavirus (strain 229E)
N:Alternate names: peplomer glycoprotein; S glycoprotein; spike glycoprotein
C:Species: human coronavirus
A:Note: host Homo sapiens (man)
C:Date: 31-Dec-1991 #sequence,revision 31-Dec-1991 #text,change 16-Jun-2000
C:Accession: A34766; S05460
R:Raabe, T.; Schelle-Prinz, B.; Siddell, S.G.

J. Gen. Virol. 71, 1065-1073, 1990
 A:Title: Nucleotide sequence of the gene encoding the spike glycoprotein of human corona
 A:Reference number: A34766; MUID:90264837
 A:Accession: A34766
 A:Molecule type: mRNA
 A:Residues: 11173 <RA>
 A:Cross-references: EMBL:X16816; NID:958926; PIDN:CAA34723.1; PID:958927
 A:Experimental source: strain 229E
 R:Raabe, T.; Sidel, S.
 Nucleic Acids Res. 17, 6387, 1989
 A:Title: Nucleotide sequence of the human coronavirus HCV 229E mRNA 4 and mRNA 5 unique
 A:Reference number: A34038; MUID:8936667
 A:Accession: S03460
 A:Status: translation not shown
 A:Molecule type: mRNA
 A:Residues: 1159-1173 <RA2>
 A:Cross-references: EMBL:X15654; NID:958921; PIDN:CAA33680.1; PID:91334827
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; transmembrane protein
 F:1-15/Domain: signal sequence #status predicted <SIG>
 F:16-1173/Product: E2 glycoprotein #status predicted <MAT>
 F:1116-1138/Domain: transmembrane #status predicted <TMN>
 F:23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,

Query Match
 Best Local Similarity 59.3%; Score 54; DB 1; Length 1173;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 KPMWPM 10
 Db 1113 KPMWPM 1119

RESULT 3
 T12505
 hypothetical protein DKFZp434C192.1 - human (fragment)
 C:Species: Homo sapiens (man)
 C>Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 23-Jul-1999
 C:Accession: T12505
 R:Ansorge, W.; Winkler, U.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
 submitted to the Protein Sequence Database, June 1999
 A:Reference number: 217527
 A:Accession: T12505
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-299 <ANS>
 A:Cross-references: EMBL:AL096753
 A:Experimental source: adult testis; clone DKFZp434C192
 C:Genetics:
 A>Note: DKFZp434C192.1

Query Match
 Best Local Similarity 86.0%; Score 51; DB 2; Length 299;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 6 PMPWPM 12
 Db 37 PMPWPM 43

RESULT 4
 B72492
 hypothetical protein APE2577 - Aeropyrum pernix (strain K1)
 C:Species: Aeropyrum pernix
 C>Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999
 C:Accession: B72492
 R:Kawarabayashi, Y.; Hino, Y.; Horikawa, H.; Yamazaki, S.; Haikawa, Y.; Jin-no, K.; Takahara, H.; Takamiya, M.; Masuda, S.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; K DNA Res. 6, 83-101, 1999
 A:Title: Complete genome sequence of an aerobic hyper-thermophilic Crenarchaeon, Aeropyrum
 A:Reference number: A72450; MUID:99310339

A:Accession: B72492
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-298 <KAW>
 A:Cross-references: DDBJ:AP000064; NID:95105945; PIDN:BAA81594.1; PID:d1045380; PID:9
 A:Experimental source: strain K1
 C:Genetics:
 A:Gene: APE2577

Query Match
 Best Local Similarity 53.8%; Score 49; DB 2; Length 298;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 2 LKPMWPM 11
 Db 102 LKPMWPM 111

RESULT 5
 J70751
 ferredoxin--NADP+ reductase (EC 1.18.1.2), long form precursor - bovine
 M:Alternate names: adrenodoxin reductase
 C:Species: Bos primigenius taurus (cattle)
 C>Date: 14-Jul-1994 #sequence_revision 18-Oct-1996 #text_change 16-Jun-2000
 C:Accession: J70751; J70079; J50390; S03558; PS0003; A29604; S52100
 R:Taketa, Y.; Sagara, Y.; Kono, A.; Sekimizu, K.; Horiochi, T.
 Biol. Pharm. Bull. 16, 1200-1206, 1993
 A:Title: Gene structure of bovine adrenodoxin reductase.
 A:Reference number: J70751; MUID:94177140
 A:Accession: J70751
 A:Molecule type: DNA
 A:Residues: 1-498 <TAK>
 A:Cross-references: GB:D83475; NID:g1199916; PIDN:BAA11921.1; PID:g4521308
 A:Experimental source: adrenal cortex
 A>Note: the authors translated the codon GTC for residue 205 as Gly
 R:Sagara, Y.; Taketa, Y.; Miyata, T.; Hara, T.; Horiochi, T.
 J. Biochem. 102, 1333-1336, 1987
 A:Title: Cloning and sequence analysis of adrenodoxin reductase cDNA from bovine adre
 A:Reference number: J70079; MUID:88198050
 A:Accession: J70079

A:Molecule type: mRNA
 A:Residues: 1-204,211-498 <SAG>
 A:Cross-references: GB:D00211; NID:g217433; PIDN:BAA00150.1; PID:g217434
 A>Note: the deduced sequence is partially confirmed by amino acid sequencing of 15 is
 R:Sagara, Y.
 submitted to DDBJ, September 1989
 A:Reference number: J50390
 A:Contents: revision, Insertion of residues 205-210
 A:Accession: J50390
 A:Molecule type: mRNA
 A:Residues: 56-498 <SAG>
 R:Hanukoglu, I.; Gutfinger, T.
 Eur. J. Biochem. 180, 479-484, 1989
 A:Title: cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites
 A:Reference number: S03558; MUID:89170752
 A:Accession: S03558
 A:Molecule type: mRNA
 A:Residues: 155-204,211-498 <HAN>
 A:Cross-references: EMBL:X13736; NID:965; PIDN:CAA32002.1; PID:g833776
 A>Note: 40S-Ser was also found
 R:Hamamoto, I.; Kurokoichi, K.; Tanaka, S.; Ichikawa, Y.
 Biochim. Biophys. Acta 953, 207-213, 1988
 A:Title: Adrenoferritin-binding peptide of NADPH-adrenoferritin reductase.
 A:Reference number: PS0003; MUID:88184054
 A:Accession: PS0003
 A:Molecule type: protein
 A:Residues: 33-41,'S',43-62,260-283,'TM',496-498 <HAM>
 A>Note: a cyanogen bromide peptide binds to adrenoferritin
 R:Nonaka, Y.; Murakami, H.; Yabuchi, Y.; Kuramitsu, S.; Kagamiyama, H.; Yamano, T.;
 Biochem. Biophys. Res. Commun. 145, 1239-1247, 1987
 A:Title: Molecular cloning and sequence analysis of full-length cDNA for mRNA of adre
 A:Reference number: A29604; MUID:87270696

A:Accession: A29604
 A:Molecule type: mRNA
 A:Residues: 1-76, 'R', 78-80, 'VMALTPRSMIL', 95-123, 'RVYPLT', 129-204, 211-273, 'R', 275-322,
 A:Cross-references: GB:M17029; NID:9162628; PIDN:AAA30362.1; PID:9162629
 A:Experimental source: adrenal cortex
 R:Wardburton, R.J.; Seybert, D.W.
 Biochim. Biophys. Acta 1246, 39-46, 1995
 A:Title: Structural and functional characterization of bovine adrenodoxin reductase by 1
 A:Reference number: S52100; MUID:95110846
 A:Accession: S52100
 A:Status: preliminary
 A:Molecule type: protein
 A:Residues: 'X', 34-41, 'X', 43-48, 'X', 50-51, 304-306, 'X', 308-309, 'X', 311-326 <MAR>
 C:Comment: Ferredoxin-NADP+ reductase is localized in the matrix of adrenal cortex mito
 ferredoxin-NADP+ reductase, adrenodoxin and two forms of cytochrome P-450.
 C:Genetics:
 A:Introns: 27/1; 59/3; 91/3; 132/3; 170/3; 204/3; 246/3; 275/1; 341/3; 399/1; 456/1
 C:Function:
 A:Description: catalyzes the reversible reduction of NADP+ by reduced ferredoxin or red
 C:Superfamily: human ferredoxin-NADP+ reductase
 C:Keywords: alternative splicing; flavoprotein; mitochondrion; monomer; NADP; oxidoreduc
 F:1-33/Domain: transit peptide (mitochondrion) #status predicted <SIG>
 F:33-498/Product: ferredoxin-NADP+ reductase, long form #status predicted <MAT>
 F:33-204, 211-498/Product: ferredoxin-NADP+ reductase, short form #status experimental <
 F:40-70/Region: beta-alpha-beta FAD nucleotide-binding fold
 F:180-190/Region: NADP binding #status predicted
 F:281/Binding site: substrate (lys) #status experimental

Query Match 53.8%; Score 49; DB 1; Length 498;
 Best Local Similarity 83.3%; Pred. No. 14;
 Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 WPMWPM 10
 I : : : :
 Db 6 WPMWPM 11

RESULT 6
 S33068
 myosin heavy chain - fluke (Schistosoma mansoni) (fragment)
 N:Alternate names: surface antigen, 200K
 C:Species: Schistosoma mansoni
 C:Date: 22-Nov-1993 #sequence_revision 06-Sep-1996 #text_change 13-Feb-1998
 C:Accession: S33068
 R:Solomon, L.M.A.; Masterson, C.P.; Tom, T.D.; McNally, M.T.; Lowell, G.H.; Strand, M.
 J. Immunol. 149, 3612-3620, 1992
 A:Title: Induction of protective immunity in mice using a 62-kDa recombinant fragment of
 A:Reference number: A46514; MUID:93056536
 A:Accession: S33068
 A:Molecule type: mRNA
 A:Residues: 1-527 <SOIT>
 A:Cross-references: EMBL:X65591
 A:Note: the authors translated the codon CAA for residue 346 as Lys
 C:Superfamily: myosin heavy chain; myosin motor domain homology
 C:Keywords: ATP; surface antigen

Query Match 53.8%; Score 49; DB 2; Length 527;
 Best Local Similarity 62.5%; Pred. No. 15;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILKKMPWM 8
 I : : : :
 Db 106 VLKMPWM 113

RESULT 7
 B70741
 probable moey protein - Mycobacterium tuberculosis (strain H37RV)
 C:Species: Mycobacterium tuberculosis
 C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 C:Accession: B70741

R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
 ; Connor, R.; Davies, R.; Devlin, K.; Feltham, T.; Gentles, S.; Hamlin, N.; Holroyd,
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 Nature 393, 537-544, 1998
 A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
 A:Reference number: A70500; MUID:98295987
 A:Accession: B70741
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-715 <COI>
 A:Cross-references: GB:Z75555; GB:AL123456; NID:93261608; PIDN:CAA99988.1; PID:e25035
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: moey

Query Match 53.8%; Score 49; DB 2; Length 715;
 Best Local Similarity 60.0%; Pred. No. 20;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 3 KKPMWPMRR 12
 I : : : :
 Db 64 KRWVYPMRR 73

RESULT 8
 A59287
 myosin heavy chain - fluke (Schistosoma mansoni) (strain Brazilian LE)
 C:Species: Schistosoma mansoni
 C:Date: 09-Jun-2000 #sequence_revision 09-Jun-2000 #text_change 08-Sep-2000
 C:Accession: A59287
 R:Weston, D.S.; Schmitz, J.; Kemp, M.; Kunz, W.
 Mol. Biochem. Parasitol. 58, 161-164, 1993
 A:Title: Cloning and sequence characterization of a complete myosin heavy chain cDNA
 A:Reference number: A59287; MUID:93211444
 A:Accession: A59287
 A:Status: preliminary; not compared with conceptual translation
 A:Molecule type: mRNA
 A:Residues: 1-1940 <WES>
 A:Cross-references: GB:L01634; PIDN:AAA29905.1
 A:Experimental source: strain Brazilian LE
 C:Genetics:
 A:Gene: MYH
 C:Superfamily: myosin heavy chain; myosin motor domain homology
 F:82-752/Domain: myosin motor domain homology <MMO>

Query Match 53.8%; Score 49; DB 2; Length 1940;
 Best Local Similarity 62.5%; Pred. No. 53;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 ILKKMPWM 8
 I : : : :
 Db 809 VLKMPWM 816

RESULT 9
 T29295
 hypothetical protein C50F7.8 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T29295
 R:Johnson, D.; Steillyes, L.
 submitted to the EMBL Data Library, November 1995
 A:Description: The sequence of C. elegans cosmid C50F7.
 A:Reference number: Z20601
 A:Accession: T29295
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-111 <COH>
 A:Cross-references: EMBL:U41557; PIDN:AAA83303.1; CESP:C50F7.8
 C:Genetics:

A:Gene: CESP:C50F7.8

Query Match
Best Local Similarity 52.7%; Score 48; DB 2; Length 111;
Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 5 PMPMPMR 12
DB 15 PMPMPGGR 22

RESULT 10

T36208
Hypothetical protein SCE36.09 - Streptomyces coelicolor

C:Species: Streptomyces coelicolor
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999

C:Accession: T36208

R:Oliver, K.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.

submitted to the EMBL Data Library, May 1999

A:Reference number: 221601

A:Accession: T36208

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1114 <COL>
A:Cross-references: EMBL:AL049763; PIDN:CAB42078.1; GSPDB:GN00070; SCOEDB:SCE36.09

A:Experimental source: strain A3(2)

A:Gene: SCOEDB:SCE36.09

Query Match
Best Local Similarity 52.2%; Score 47.5; DB 2; Length 114;
Matches 7; Conservative 2; Mismatches 1; Indels 1; Gaps 1;

OY 3 KKM-PMPMPMR 12
DB 102 RMRPMPMR 112

RESULT 11

S23449
NADH oxidase (H2O2-forming) (EC 1.6.-.-) - Thermus aquaticus

C:Species: Thermus aquaticus
C:Date: 22-Jan-1993 #sequence_revision 22-Jan-1993 #text_change 23-Mar-1993

C:Accession: S23449; S24556

R:Park, H.J.; Kreutzer, R.; Reiser, C.O.A.; Sprinzl, M.

Eur. J. Biochem. 205, 875-879, 1992

A:Title: Molecular cloning and nucleotide sequence of the gene encoding a H(2)O(2)-form

A:Reference number: S23449; MUID:92249331

A:Accession: S23449

A:Molecule type: DNA

A:Residues: 1-248 <PAR>

A:Cross-references: EMBL:X60110

A:Accession: S24556

A:Molecule type: protein

A:Residues: 1-32 <PAR1>

C:Genetics:

A:Gene: nox

C:Keywords: NAD; oxidoreductase

F:1-248/Product: NADH oxidase (H2O2-forming) #status experimental <MAT>

Query Match
Best Local Similarity 51.6%; Score 47; DB 2; Length 248;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 PMPMP 10
DB 179 PMPMP 183

RESULT 12

G70715
Hypothetical protein RV0945 - Mycobacterium tuberculosis (strain H37RV)

C:Species: Mycobacterium tuberculosis

C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 20-Jun-2000

C:Accession: G70715

R:Coyle, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon

; Connor, R.; Davies, R.; Felwell, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,

Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skellon, S.; Squares, S.

Nature 393, 537-544, 1998

A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno

A:Reference number: A70500; MUID:98295987

A:Accession: G70715

A:Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1253 <COL>

A:Cross-references: GB:T279700; GB:AL123456; NID:g3261628; PIDN:CAB02005.1; PID:g15242

A:Experimental source: strain H37RV

C:Genetics:

A:Gene: RV0945

C:Superfamily: ribitol dehydrogenase; short-chain alcohol dehydrogenase homology

F:8-190/Domain: short-chain alcohol dehydrogenase homology <SADH>

Query Match
Best Local Similarity 51.6%; Score 47; DB 2; Length 253;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 PMPMP 10
DB 230 PMPMP 234

RESULT 13

S77334
histidinol-phosphate aminotransferase hsc-1 - Synechocystis sp. (strain PCC 6803)

N:Alternate names: protein sl11713

C:Species: Synechocystis sp.

A:Variety: PCC 6803

C:Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 20-Jun-2000

C:Accession: S77334

R:Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima,

O. K.; Okumura, S.; Shingo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yas

DNA Res. 3, 109-136, 1996

A:Title: Sequence analysis of the genome of the unicellular cyanobacterium Synechocys

A:Reference number: S74322; MUID:97061201

A:Accession: S77334

A:Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-352 <KAN>

A:Cross-references: EMBL:D90906; GB:AE001339; NID:g1652492; PIDN:BAI17457.1; PID:g165

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

C:Genetics:

A:Gene: hsc-1

C:Superfamily: probable histidinol-phosphate transaminase

Query Match
Best Local Similarity 51.1%; Score 46.5; DB 2; Length 352;
Matches 8; Conservative 1; Mismatches 2; Indels 5; Gaps 1;

OY 2 LKKMPW-----WPMR 12
DB 106 LKTIWQVDQWPMR 121

RESULT 14

S37664
peplomer protein precursor - avian infectious bronchitis virus (strain D1466) (

N:Contains: E2 glycoprotein subunit S2

C:Species: avian infectious bronchitis virus, IBV

A:Variety: strain D1466

C>Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 20-Sep-1999
 C:Accession: S37664
 R:Kusters, J.G.; Jager, E.J.; Niesters, H.G.M.; van der Zeijst, B.A.M.
 Vaccine 8, 605-608, 1990
 A>Title: Sequence evidence for RNA recombination in field isolates of avian coronavirus
 A:Reference number: S37663; MUID:91205880
 A:Accession: S37664
 A:Molecule type: genomic RNA
 A:Residues: 1-621 <KUS>
 A:Cross-references: EMBL:X58001; NID:g58986; PIDN:CAA41065.1; PID:g58987
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; peplomer protein; spike protein
 F:1-5/Product: E2 glycoprotein subunit S1 (fragment) #status predicted <GS1>
 F:6-621/Product: E2 glycoprotein subunit S2 #status predicted <GS2>
 F:10,47,59,137,144,415,447,482,506,519,542/Binding site: carbohydrate (Asn) (covalent) #

Query Match 51.1%; Score 46.5; DB 2; Length 621;
 Best Local Similarity 61.5%; Pred. No. 38;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;
 QY 1 ILK--KMPMPW 10
 ||| ||||: |
 Db 553 ILKTYIKMPWYVW 565

RESULT 15
 S37663
 Peplomeric polyprotein precursor - avian infectious bronchitis virus (strain D207) (frag
 N:Contains: E2 glycoprotein subunit S2
 C:Species: avian infectious bronchitis virus, IBV
 A:Variety: strain D207
 C>Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 20-Sep-1999
 C:Accession: S37663
 R:Kusters, J.G.; Jager, E.J.; Niesters, H.G.M.; van der Zeijst, B.A.M.
 Vaccine 8, 605-608, 1990
 A>Title: Sequence evidence for RNA recombination in field isolates of avian coronavirus
 A:Reference number: S37663; MUID:91205880
 A:Accession: S37663
 A:Molecule type: genomic RNA
 A:Residues: 1-630 <KUS>
 A:Cross-references: EMBL:X58003; NID:g58988; PIDN:CAA41067.1; PID:g58989
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; peplomer protein; spike protein
 F:1-5/Product: E2 glycoprotein subunit S1 (fragment) #status predicted <GS1>
 F:6-621/Product: E2 glycoprotein subunit S2 #status predicted <GS2>
 F:10,47,59,137,144,415,447,482,506,519,542/Binding site: carbohydrate (Asn) (covalent) #

Query Match 51.1%; Score 46.5; DB 2; Length 630;
 Best Local Similarity 61.5%; Pred. No. 39;
 Matches 8; Conservative 1; Mismatches 1; Indels 3; Gaps 1;

QY 1 ILK--KMPMPW 10
 ||| ||||: |
 Db 553 ILKTYIKMPWYVW 565

Search completed: January 4, 2002, 08:41:32
 Job time: 170 sec

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AUTHORS Sasaki, T. and Yamamoto, K.
 TITLE Rice cDNA from green shoot (2001)
 JOURNAL Unpublished (2001)
 COMMENT Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasaki@agr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
 PROJECT = 'RGP',
 S16019_97A.

FEATURES

source Location/Qualifiers

1. 448
 /organism="Oryza sativa"
 /strain="Nipponbare"
 /db_xref="taxon:4530"
 /clone="S16019"
 /clone_lib="Rice green shoot"
 /note="Green shoot (8 days old)"
 BASE COUNT 85 a 146 c 160 g 57 t
 ORIGIN

alignment_scores:
 Quality: 69.00 Length: 10
 Ratio: 7.667 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-36 x AU198162/rev ..

Align seg 1/1 to reverse of: AU198162 from: 1 to: 448

3 ArgTrrpTrpTrpProTrrpArgArglys 12
 |||||
 311 CGCTGGCCTTGCTGGCCCTGACCGCGCG 282

seq_name: gb_est1:AU082117

seq_documentation_block:
 LOCUS AU082117 578 bp mRNA EST 04-FEB-2000
 DEFINITION AU082117 Rice panicle at ripening stage Oryza sativa cDNA clone
 E11611, mRNA sequence.
 ACCESSION AU082117
 VERSION AU082117.1 GI:6727452
 KEYWORDS EST.
 SOURCE Oryza sativa.
 ORGANISM Oryza sativa
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 Eriatidae; Oryzoideae; Oryzae; Oryza.
 1 (bases 1 to 578)

REFERENCE
 AUTHORS Sasaki, T. and Yamamoto, K.
 TITLE Rice cDNA from panicle at ripening stage (2000)
 JOURNAL Unpublished (2000)
 COMMENT Contact: Takuji Sasaki
 National Institute of Agrobiological Resources
 Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
 305-8602, Japan
 Tel: 81-298-38-7441
 Fax: 81-298-38-7468
 Email: tsasaki@agr.affrc.go.jp, URL: http://rgp.dna.affrc.go.jp/
 PROJECT = 'RGP',

FEATURES

source Location/Qualifiers

1. 578
 /organism="Oryza sativa"
 /strain="Nipponbare"
 /db_xref="taxon:4530"
 /clone="E11611"
 /clone_lib="Rice panicle at ripening stage"
 /dev_stage="ripening stage"
 /note="Organ: panicle; Rice cDNA from panicle at ripening

BASE COUNT 129 a 180 c 179 g 90 t
 ORIGIN

alignment_scores:
 Quality: 69.00 Length: 10
 Ratio: 7.667 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-09-444-281-36 x AU082117/rev ..

Align seg 1/1 to reverse of: AU082117 from: 1 to: 578

3 ArgTrrpTrpTrpProTrrpArgArglys 12
 |||||
 345 CGCTGGCCTTGCTGGCCCTGACCGCGCG 316

seq_name: gb_est1:AU089922

seq_documentation_block:

LOCUS AU089922 446 bp mRNA EST 19-APR-2000
 DEFINITION AU089922 Hordeum vulgare subsp. vulgare upper three leaves at
 heading stage Hordeum vulgare subsp. vulgare cDNA clone
 haruna_1lib1_121, mRNA sequence.
 haruna_1lib1_121, mRNA sequence.
 ACCESSION AU089922
 VERSION AU089922.1 GI:7613350
 KEYWORDS EST.
 SOURCE Hordeum vulgare subsp. vulgare.
 ORGANISM Hordeum vulgare subsp. vulgare
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae
 ; Triticeae; Hordeum.
 1 (bases 1 to 446)

REFERENCE
 AUTHORS Sato, K., Takahashi, H. and Takeda, K.
 TITLE Hordeum vulgare subsp. vulgare cDNA clone
 JOURNAL Unpublished (2000)
 COMMENT Contact: Kazuhiko Sato
 Research Institute for Bioreources
 Okayama University, Barley Germplasm Center
 Chuo 2-20-1, Kurashiki, Okayama 710-0046, Japan
 Email: kazsato@rib.okayama-u.ac.jp
 URL: http://www.rib.okayama-u.ac.jp/barley/.

FEATURES

source Location/Qualifiers

1. 446
 /organism="Hordeum vulgare subsp. vulgare"
 /cultivar="Haruna N130"
 /db_xref="taxon:112509"
 /clone="haruna_1lib1_121"
 /clone_lib="Hordeum vulgare subsp. vulgare upper three
 leaves at heading stage"
 /tissue_type="Upper three leaves at heading stage"
 BASE COUNT 89 a 130 c 149 g 76 t 2 others
 ORIGIN

alignment_scores:
 Quality: 68.00 Length: 8
 Ratio: 8.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-09-444-281-36 x AU089922/rev ..

Align seg 1/1 to reverse of: AU089922 from: 1 to: 446

4 TrpProTrrpTrpProTrrpArgArg 11
 |||||
 188 TGGCGTGCTGGCCGCGCGCGA 165

seq_name: gb_est1:AU089934


```

Durham, NC 27708-1000, USA
Tel: 919 613 8164
Fax: 919 613 8177
Email: chlamy@duke.edu
location/Qualifiers
1..330
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type mt+ 21gr"
/db_xref="taxon:3055"
/cclone_lid="C. reinhardtii CC-1690, normalized, Lambda Zap
I1"
/notes="Vector: pBluescript II SK-; Site.1: EcoRI; Site.2:
XhoI; This library, constructed by John Davies and Jeffrey
Middermott, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP (acetate-containing) medium in the
light, TAP medium in the dark, HS (minimal) medium in
ambient levels of CO2 and HS medium bubbled with 5% CO2.
POLYA mRNA was purified from each sample, pooled and cDNA
synthesized. The cDNA was directionally cloned into lambda
ZAP II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
pBluescript II SK- plasmids were excised from the lambda
ZAP clones by superinfection with Exassist (Stratagene)
phase. The library was normalized using method 4 described
in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT      40 a      103 c      142 g      45 t
ORIGIN

alignment_scores:
    Quality:      67.00      Length:      9
    Ratio:        8.375      Gaps:        0
    Percent Similarity: 88.889      Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x BE024584 ..

Align seg 1/1 to: BE024584 from: 1 to: 350

3 ArgTrpProTrpTrpProTrpArg 11
|||||
72 CGGTGGCGGCGGTGGCGGCGGCGG 98

seq_name: gb_est1:A0198258

seq_documentation_block:
LOCUS      A0198258      352 bp      mRNA      EST      12-JUL-2001
DEFINITION A0198258 Rice green shoot Oryza sativa cDNA clone S16389, mRNA
sequence.
ACCESSION A0198258
VERSION A0198258.1 GI:14714335
KEYWORDS EST.
SOURCE Oryza sativa.
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
Ehartoideae; Oryzaceae; Oryza.
1 (bases 1 to 352)
Sasaki,T. and Yamamoto,K.
Rice cDNA from green shoot (2001)
Unpublished (2001)
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program, Kannondai 2-1-2, Tsukuba, Ibaraki
305-8602, Japan
Tel: 81-298-38-7441
Fax: 81-298-38-7468
Email: tsasaki@affrc.go.jp, URL:http://rgp.dna.affrc.go.jp/
PROJECT = 'RGP'.
S16389_96Z.

FEATURES
Source
Location/Qualifiers
1..352
/organism="Oryza sativa"
/strain="Nipponbare"

```

/db_xref="taxon:4530"
/clone="S16389"
/clone_lib="Rice green shoot"
/note="Green shoot (8 days old)"
BASE COUNT 95 a 95 c 104 g 51 t 7 others
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AU198258/rev ..

Align seg 1/1 to reverse of: AU198258 from: 1 to: 352

3 ArgTrpProTrrPrrProTrrPArgArg 11
|||||
167 CGCTGGCGCTTGCTGGCCCTGACGCGG 141

seq_name: gb_est1:AV641634

seq_documentation_block:
LOCUS AV641634 371 bp mRNA EST 15-DEC-2000
DEFINITION AV641634 Chlamydomonas reinhardtii 5% CO2 Chlamydomonas reinhardtii
ACCESSION AV641634
VERSION AV641634.1 GI:10784962
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.

REFERENCE 1 (bases 1 to 371)
Asanuma,E., Miura,K., Inoue,Y., Fukuzawa,H., Ohyama,K.,
Nakamura,Y. and Tabata,S.
Generation of expressed sequence tags from low-CO2 and high-CO2
adapted cells of Chlamydomonas reinhardtii
DNA Res. 7 (5), 405-307 (2000)

JOURNAL MEDLINE
20539644
Contact: Erika Asanuma
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: asanumakazusa.or.jp. URL: http://www.kazusa.or.jp/en/plant/.

FEATURES
source
1..371
Location/Qualifiers
/organism="Chlamydomonas reinhardtii"
/strain="C9"
/db_xref="taxon:3055"
/clone_lib="HCL037H03_F"
/note="Vector: Bluescript II SK-; Site_1: EcoRI; Site_2:
XhoI; The cDNA library was constructed from cells cultured
in a medium with bubbling air containing 5% carbon
dioxide"

BASE COUNT 73 a 106 c 141 g 51 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x AV641634 ..

Align seg 1/1 to: AV641634 from: 1 to: 371

3 ArgTrpProTrrPrrProTrrPArgArg 11

|||||
244 CGGTGGCGCTTGCTGGCCCTGACGCGG 270

seq_name: gb_est1:BE129188

seq_documentation_block:
LOCUS BE129188 386 bp mRNA EST 21-JUN-2000
DEFINITION 894021E12.Y1 C. reinhardtii CC-1690, normalized, Lambda Zap II
Chlamydomonas reinhardtii cDNA, mRNA sequence.

ACCESSION BE129188
VERSION BE129188.1 GI:8576551
KEYWORDS EST.
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.

REFERENCE 1 (bases 1 to 386)
Grossman,A., Davies,J., Federspiel,N., Harris,E., Lefebvre,P.,
McDermott,J.P., Sillflow,C., Stern,D. and Surzycki,R.
Analyses of the Chlamydomonas reinhardtii Genome: A Model,
Unicellular System for Analyzing Gene Function and Regulation in
Vascular Plants: project phase 2

JOURNAL

COMMENT
Unpublished (2000)
Contact: Elizabeth H. Harris
DCMB Box 91000
Duke University
Durham, NC 27708-1000, USA
Tel: 919 613 8164
Fax: 919 613 8177
Email: chlamyduke.edu.

FEATURES

source
1..386
Location/Qualifiers
/organism="Chlamydomonas reinhardtii"
/strain="CC-1690 wild type mt+ 21gr"
/db_xref="taxon:3055"
/clone_lib="C. reinhardtii CC-1690, normalized, Lambda Zap
II"

/note="Vector: Bluescript II SK-; Site_1: EcoRI; Site_2:
XhoI; This library, constructed by John Davies and Jeffrey
McDermott, combines cDNAs from CC-1690 cells grown to
mid-log phase in TAP (acetate-containing) medium in the
light, TAP medium in the dark, HS (minimal) medium in
ambient levels of CO2 and HS medium bubbled with 5% CO2.
Polya mRNA was purified from each sample, pooled and cDNA
synthesized. The cDNA was directionally cloned into lambda
ZAP II (Stratagene) in the EcoRI (5') and XhoI (3') sites.
Bluescript II SK- plasmids were excised from the lambda
ZAP clones by superinfection with EXAssist (Stratagene)
phage. The library was normalized using method 4 described
in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT 79 a 109 c 142 g 56 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 9
Ratio: 8.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x BE129188 ..

Align seg 1/1 to: BE129188 from: 1 to: 386

3 ArgTrpProTrrPrrProTrrPArgArg 11
|||||
249 CGGTGGCGCTTGCTGGCCCTGACGCGG 275

seq_name: gb_est2:BF584442

seq_documentation_block:
LOCUS BF584442 701 bp mRNA EST 12-DEC-2000
DEFINITION 602098336F1 NCL_CGAP_Co24 Mus musculus cDNA clone IMAGE:4218273 5',

ACCESSION mRNA sequence.
BF584442
VERSION BF584442.1 GI:11658160
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 701)
TITLE NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
Plate: LLM9798 row: g column: 10
High quality sequence stop: 701.

FEATURES
source
1. 701
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone_image="4218273"
/lab_host="NCI_CGAP_C024"
/note="Organ: colon; Vector: PCMV-SpOrf; Site: 1: NotI;
Site: 2: SalI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.6 kb. Constructed by Life
Technologies. Note: this is a NCI_CGAP library."

BASE COUNT 149 a 198 c 186 g 168 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 10
Ratio: 7.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x BF584442
Align seg 1/1 to: BF584442 from: 1 to: 701

11leuAgtTrrpTrpTrpTrpArg 10
:::||||| ||||||| ||||||| |||||||
188 CTGTATTTGTGCGCTTGTGCGCATGAGCA 217

seq_name: gb_est2:BG964576

seq_documentation_block:
LOCUS BG964576 834 bp mRNA EST 12-JUN-2001
DEFINITION 6028332551 NCI_CGAP_C024 Mus musculus cDNA clone IMAGE:4986828 5',
mRNA sequence.
ACCESSION BG964576
VERSION BG964576.1 GI:14352213
KEYWORDS EST.
SOURCE house mouse.
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 834)
TITLE NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs@mail.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.

CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov
Plate: LLM10996 row: n column: 13
High quality sequence stop: 811.

FEATURES
source
1. 834
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone_image="4986828"
/lab_host="NCI_CGAP_C024"
/note="Organ: colon; Vector: PCMV-SpOrf; Site: 1: NotI;
Site: 2: SalI; Cloned unidirectionally. Primer: Oligo dT.
Average insert size 1.6 kb. Constructed by Life
Technologies. Note: this is a NCI_CGAP library."

BASE COUNT 185 a 202 c 226 g 221 t
ORIGIN

alignment_scores:
Quality: 67.00 Length: 10
Ratio: 7.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x BG964576
Align seg 1/1 to: BG964576 from: 1 to: 834

11leuAgtTrrpTrpTrpTrpArg 10
:::||||| ||||||| ||||||| |||||||
195 CTGTATTTGTGCGCTTGTGCGCATGAGCA 224

seq_name: gb_gss:A0943724

seq_documentation_block:
LOCUS A0943724 514 bp DNA GSS 27-JAN-2000
DEFINITION Sheared DNA-36H6.TF Sheared DNA Trypanosoma brucei genomic clone
Sheared DNA-36H6, DNA sequence.
ACCESSION A0943724
VERSION A0943724.1 GI:6766989
KEYWORDS GSS.
SOURCE Trypanosoma brucei.
ORGANISM Trypanosoma brucei
REFERENCE Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 514)
TITLE El-Sayed, N., Zhao, S., Zhao, H., Gill, S., Suh, E., Malek, J., Fujii, C.,
Gerrard, C., Leech, V., de Jong, P., Ullu, E., Melville, S., Donelson, J.,
Fraser, C. and Adams, M.
Determination of clone end sequences from Trypanosoma brucei cDNA
10.1 sheared DNA library
Unpublished (1999)
Other_GSSs: Sheared DNA-36H6.TF
Contact: Najib M. El-Sayed
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: nelsayed@tigr.org
Clones are derived from the Trypanosoma brucei cDNA 10.1 sheared
DNA library constructed at TIGR. Clones will be available for
distribution through ATCC. Sheared DNA end sequences search page:
http://www.tigr.org/tdb/mdb/cdbd/
Seq primer: M13-Forward
Class: shotgun.
Location/Qualifiers

source
1. 514
/organism="Trypanosoma brucei"
/strain="TREU927/4 GUTat 10.1"
/db_xref="taxon:5691"
/clone="Sheared DNA-36H6"
/note="Vector: pUC18; Site_1: Small; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrell, Oxford University Press, 1999)."

BASE COUNT 124 a 185 c 131 g 74 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-09-444-281-36 x AQ943724/rev ..
Align seg 1/1 to reverse of: AQ943724 from: 1 to: 514

2 LeuArgTTProtTTProtTTProtTTPargArg 11
|||||
450 CTTCGATGGCCTTGCTGTGTGTGGTGGCGCG 421

seq_name: gb_gss:A2218599
seq_documentation_block:
LOCUS A2218599 572 bp DNA GSS 09-JUN-2000
DEFINITION Sheared DNA-82B1.TR Sheared DNA Trypanosoma brucei genomic clone
ACCESSION A2218599
VERSION A2218599.1 GI:8436399
KEYWORDS GSS.
SOURCE Trypanosoma brucei.
ORGANISM Trypanosoma brucei.
Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae;
Trypanosoma.
1 (bases 1 to 572)
El-Sayed,N., Zhao,S., Zhao,H., Gill,S., Suh,E., Malek,J., Fujii,C.,
Gerard,C., Leech,V., de Jong,P., Ullu,E., Melville,S., Donelson,J.,
Fraser,C. and Adams,M.
Determination of clone end sequences from Trypanosoma brucei GUTat
10.1 sheared DNA library
Unpublished (1999)
Other_GSSs: Sheared DNA-82B1.TF
Contact: Najib M. El-Sayed
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: nelsayed@tigr.org
Clones are derived from the Trypanosoma brucei GUTat 10.1 sheared
DNA library constructed at TIGR. Clones will be available for
distribution through Research Genetics, Alabama, USA. Sheared DNA
end sequences search page: <http://www.tigr.org/tdb/mbd/tbdt/>.
Seq primer: M13-Reverse
Class: shotgun.
FEATURES
location/Qualifiers
1..572
/organism="Trypanosoma brucei"
/strain="TREU927/4 GUTat 10.1"

/db_xref="taxon:5691"
/clone="Sheared DNA-82B1"
/clone_lib="Sheared DNA"
/note="Vector: pUC18; Site_1: Small; Constructed at The Institute for Genomic Research (TIGR), Rockville, MD. Genomic DNA isolated from a cloned population of Trypanosoma brucei (TREU927/4 GUTat 10.1) was mechanically sheared to give a tight size distribution (approx 2 kb). The v + i method used for the library construction is described in detail in Smith, H.O. and Venter, J.C. (Making small insert libraries for whole genome shotgun sequencing projects. In Genome Sequencing: A Practical Approach, eds. M. Vaubin and B. Barrell, Oxford University Press, 1999)."

BASE COUNT 136 a 217 c 144 g 75 t
ORIGIN

alignment_scores:
Quality: 66.00 Length: 10
Ratio: 7.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-09-444-281-36 x A2218599/rev ..
Align seg 1/1 to reverse of: A2218599 from: 1 to: 572

2 LeuArgTTProtTTProtTTProtTTPargArg 11
|||||
211 CTTCGATGGCCTTGCTGTGTGTGGTGGCGCG 182

seq_name: gb_est:C52835
seq_documentation_block:
LOCUS C52835 208 bp mRNA EST 11-SEP-1997
DEFINITION C52835 yuji Kohara unpublished cdna Caenorhabditis elegans CDNA
clone yk285b1 3', mRNA sequence.
ACCESSION C52835
VERSION C52835.1 GI:2390592
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea
; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 208)
Kohara,Y., Morishashi,T., Tabara,H., Watanabe,H., Sugimoto,A., Sano
M., Miyata,A. and Nishigaki,A.
Expression map of the C.elegans genome
Unpublished (1996)
Contact: Yuji Kohara
Genome Biology Lab.
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
location/Qualifiers
1..208
/organism="Caenorhabditis elegans"
/strain="CHI489 him-8(e1489)"
/db_xref="taxon:6239"
/clone="yk285b1"
/clone_lib="yuji Kohara unpublished cdna"
/sex="hermaphrodite, male"
/tissue_type="whole animal"
/dev_stage="varied"

BASE COUNT 25 a 50 c 77 g 52 t 4 others
ORIGIN

alignment_scores:
Quality: 65.00 Length: 10

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GenCore version 4.5
Copyright (c) 1993 - 2000 CompuGen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:24 ; Search time 50.17 Seconds
(Without alignments)
34.986 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRPMWPMRRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 473505 seqs, 146272329 residues
Total number of hits satisfying chosen parameters: 473505

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: SPTEMBL_17:*
2: sp_archaea:*
3: sp_bacteria:*
4: sp_fungi:*
5: sp_human:*
6: sp_invertebrate:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	57	66.3	723	12 Q9DUC4	Q9DUC4 tt virus. o
2	53	61.6	746	12 Q9JH31	Q9JH31 tt virus. o
3	53	61.6	1173	12 Q990M4	Q990M4 human coron
4	53	61.6	1173	12 Q990M3	Q990M3 human coron
5	53	61.6	1173	12 Q990M2	Q990M2 human coron
6	53	61.6	1173	12 Q990M1	Q990M1 human coron
7	53	61.6	1383	12 Q84712	Q84712 porcine epi
8	52	60.5	1245	3 Q9Y7V5	Q9Y7V5 trichodema
9	51	59.3	299	4 Q9YVNI	Q9YVNI homo sapien
10	51	59.3	504	2 P96143	P96143 thermococin
11	50	58.1	141	5 Q18753	Q18753 caenorhabd
12	50	58.1	141	5 Q9CZAI	Q9CZAI mus musculu
13	50	58.1	327	10 Q9AUN3	Q9AUN3 oryza sativ
14	50	58.1	735	12 Q9DUC9	Q9DUC9 tt virus. o
15	49	57.0	49	12 Q9D80	Q9D80 tt virus. o
16	49	57.0	226	4 Q9BS58	Q9BS58 homo sapien
17	49	57.0	467	5 Q19573	Q19573 caenorhabd
18	49	57.0	748	12 Q9D81	Q9D81 tt virus. o
19	48.5	56.4	114	2 Q9X8C2	Q9X8C2 streptomyce

20	48	55.8	540	2 Q07504	Q07504 bacillus me
21	47	54.7	165	10 Q9SNN3	Q9SNN3 oryza sativ
22	47	54.7	276	2 Q9HXC9	Q9HXC9 pseudomonas
23	47	54.7	1411	10 Q9LYG0	Q9LYG0 arabidopsis
24	46	53.5	154	2 Q9R6J3	Q9R6J3 agrobacteri
25	46	53.5	728	3 Q9P3G0	Q9P3G0 neurospora
26	45	52.3	159	2 Q9KZT3	Q9KZT3 streptomyce
27	45	52.3	273	2 Q9PCR3	Q9PCR3 xyloella fas
28	45	52.3	412	2 Q916P7	Q916P7 pseudomonas
29	45	52.3	423	2 Q24742	Q24742 bacteroides
30	45	52.3	443	10 Q9S751	Q9S751 oryza sativ
31	45	52.3	448	2 Q9WYR8	Q9WYR8 thermotoga
32	45	52.3	525	10 Q9ATU5	Q9ATU5 lolium rigi
33	45	52.3	525	10 Q9ATU2	Q9ATU2 lolium rigi
34	45	52.3	525	10 Q9ATU1	Q9ATU1 lolium rigi
35	45	52.3	730	10 Q9RG26	Q9RG26 arabidopsis
36	45	52.3	767	12 Q9QUD8	Q9QUD8 tt virus. h
37	45	52.3	1100	11 Q921J9	Q921J9 mus musculu
38	44.5	51.7	766	12 Q91FV0	Q91FV0 tt virus. p
39	44	51.2	143	4 Q9H9A4	Q9H9A4 homo sapien
40	44	51.2	145	2 Q86437	Q86437 pseudomonas
41	44	51.2	257	2 Q56924	Q56924 yersinia en
42	44	51.2	361	2 Q9P1W6	Q9P1W6 homo sapien
43	44	51.2	406	5 Q9W404	Q9W404 drosophila
44	44	51.2	429	5 Q9N8Y2	Q9N8Y2 trypanosoma
45	44	51.2	458	4 Q9UCB1	Q9UCB1 homo sapien

ALIGNMENTS

RESULT 1
ID Q9DUC4 PRELIMINARY: PRT: 723 AA.
AC Q9DUC4;
DT 01-MAR-2001 (TREMblrel. 16, Created)
DT 01-MAR-2001 (TREMblrel. 16, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE ORF1.
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_Taxid=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RA Okamoto H.;
RL Submitted (APR-2000) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=MF-TTV9;
RX PubMed=11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TT viruses in humans and nonhuman primates and their
RT phylogenetic relatedness";
RL Virology 277:368-378(2000).
DR EMBL: AB041959; BAB19313.1;
DR InterPro: IPR001563; Serine-carboxypept.
DR PROSITE: PS00131; CARBOXYPEPT_SER; UNKNOWN.1.
SQ SEQUENCE 723 AA; 85393 MW; 232D003098766344 CRC64;

Query Match 66.3%; Score 57; DB 12; Length 723;
Best Local Similarity 100.0%; Pred. No. 4.0;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 5 PMWPMRR 11
Db 2 PMWPMRR 8
RESULT 2
Q9JH31

ID 09JH31 PRELIMINARY; PRT; 746 AA.
AC 09JH31;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DE 01-OCT-2000 (TREMBlrel. 15, Last annotation update)
DE 09JH31;
OS Viruses; ssDNA viruses; unclassified ssDNA viruses.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
NCBI_TaxID=68887;
RN 11;
RP SEQUENCE FROM N.A.
RC STRAIN=TUN02;
RA Okamoto H.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN 12;
RP SEQUENCE FROM N.A.
RC STRAIN=TUN02;
RA Ukita M., Okamoto H., Nishizawa T., Tawara A., Takahashi M.,
RA Iizuka H., Miyakawa Y., Mayumi M.;
RT "The entire nucleotide sequences of two distinct TT virus (TTV)
RT isolates (TUN01 and TUN02) remotely related to the original TTV
RT isolates.";
RL Arch. Virol. 0:0-0(2000).
DR EMBL: AB028669; BAA94878.1; -
SQ SEQUENCE 746 AA; 88561 MW; E0B22953AE764E3E CRC64;

Query Match 61.6%; Score 53; DB 12; Length 746;
Best Local Similarity 54.5%; Pred. No. 15;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 2 LRPMWPMWRK 12
Db 1 MANGWWRWRRR 11

RESULT 3
0990M4 PRELIMINARY; PRT; 1173 AA.
ID 0990M4;
AC 0990M4;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronavirusidae; Coronavirus.
OX NCBI_TaxID=11137;
RN 11;
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected
RT with human coronavirus HCoV-229E.";
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344186; AAK32188.1; -
SQ SEQUENCE 1173 AA; 128669 MW; ABC6E0A75E8BD8A4 CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
Best Local Similarity 62.5%; Pred. No. 23;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWPMW 9
Db 1112 IKPMWVW 1119

RESULT 4
0990M3 PRELIMINARY; PRT; 1173 AA.
ID 0990M3;
AC 0990M3;

DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronavirusidae; Coronavirus.
OX NCBI_TaxID=11137;
RN 11;
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected
RT with human coronavirus HCoV-229E.";
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344187; AAK32189.1; -
SQ SEQUENCE 1173 AA; 128683 MW; 9E2368160082A81A CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
Best Local Similarity 62.5%; Pred. No. 23;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWPMW 9
Db 1112 IKPMWVW 1119

RESULT 5
0990M2 PRELIMINARY; PRT; 1173 AA.
ID 0990M2;
AC 0990M2;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronavirusidae; Coronavirus.
OX NCBI_TaxID=11137;
RN 11;
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected
RT with human coronavirus HCoV-229E.";
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344188; AAK32190.1; -
SQ SEQUENCE 1173 AA; 128653 MW; 8B658FCBBD1842DA CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1173;
Best Local Similarity 62.5%; Pred. No. 23;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRPMWPMW 9
Db 1112 IKPMWVW 1119

RESULT 6
0990M1 PRELIMINARY; PRT; 1173 AA.
ID 0990M1;
AC 0990M1;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE GLYCOPROTEIN.
GN S.
OS Human coronavirus (strain 229E).
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;

OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=1137;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=229E;
RA Bonavia A., Holmes K.V.;
RT "Viral and cellular changes in a human cell line persistently infected
with human coronavirus HCoV-229E."
RL Submitted (FEB-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF344189; AAK32191.1; -
SQ SEQUENCE 1173 AA; 128760 MW; B73A165A6270152A CRC64;

Query Match
Best Local Similarity 61.6%; Score 53; DB 12; Length 1173;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRMPMPW 9
Db 1112 IKMPMWV 1119

RESULT 7
ID 084712 PRELIMINARY; PRT; 1383 AA.
AC 084712;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-JUN-2001 (TREMBlrel. 17, Last annotation update)
DE SPIKE PROTEIN.
OS porcine epidemic diarrhea virus.
OC Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;
OC Coronaviridae; Coronavirus.
OX NCBI_TaxID=28295;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=94231173; PubMed=8176382;
RA Duarte M., Laude H.;
RT "Sequence of the spike protein of the porcine epidemic diarrhoea
virus."
RL J. Gen. Virol. 75:1195-1200(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=93389433; PubMed=8397280;
RA Bridgen A., Duarte M., Tobler K., Laude H., Ackermann M.;
RT "Sequence determination of the nucleocapsid protein gene of the
porcine epidemic diarrhoea virus confirms that this virus is a
coronavirus related to human coronavirus 229E and porcine
transmissible gastroenteritis virus."
RL J. Gen. Virol. 74:1795-1804(1993).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=BRI/87;
RX MEDLINE=94120721; PubMed=8291230;
RA Duarte M., Tobler K., Bridgen A., Rasschaert D., Ackermann M.,
Laude H.;
RT "Sequence analysis of the porcine epidemic diarrhoea virus genome
between the nucleocapsid and spike protein genes reveals a polymorphic
ORF."
RL Virology 198:466-476(1994).
DR EMBL: Z25483; CAA80971.1; -
DR InterPro: IPR002551; Corona_S1.
DR InterPro: IPR002552; Corona_S2.
DR Pfam: PF01600; Corona_S1; 1.
DR Pfam: PF01601; Corona_S2; 1.
FT CONFLICT 422 422 Y -> N (IN REF. 1).
SQ SEQUENCE 1383 AA; 151404 MW; 741C84D5DD3BDC4D CRC64;

Query Match 61.6%; Score 53; DB 12; Length 1383;
Best Local Similarity 62.5%; Pred. No. 27;

Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
OY 2 LRMPMPW 9
Db 1321 IKMPMWV 1328

RESULT 8
ID 0977V5 PRELIMINARY; PRT; 1245 AA.
AC 0977V5;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE CONIDIOSPORE SURFACE PROTEIN.
GN CMPI.
OS Trichoderma harzianum.
OC Eukaryota; Fungi; Ascomycota; mitosporic Ascomycota; Trichoderma.
OX NCBI_TaxID=5544;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 32173;
RA Puyesky M., Benhamou N., Ponce Noyola P., Bauw G., Ziv T.,
van Montagu M., Herrera Estrella A., Horwitz B.A.;
RT "Developmental regulation of a gene encoding a multidomain
RT conidiospore surface protein of Trichoderma, cmpl."
RL Submitted (MAR-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AJ133651; CAB40845.1; -
SQ SEQUENCE 1245 AA; 135824 MW; 3249C749AFA0CDF8 CRC64;

Query Match 60.5%; Score 52; DB 3; Length 1245;
Best Local Similarity 60.0%; Pred. No. 33;
Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

OY 3 RMPMPMPRRK 12
Db 1185 RMQWMSWPRR 1194

RESULT 9
ID 09Y4N1 PRELIMINARY; PRT; 299 AA.
AC 09Y4N1;
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAY-2000 (TREMBlrel. 13, Last annotation update)
DE HYPOTHETICAL 34.0 KDA PROTEIN (FRAGMENT).
GN DKFZP434C192.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=TESTIS;
RA Ansoorge W., Winkner U., Mewes H.W., Gassenhuber J., Wiemann S.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL096753; CAB46428.2; -
KW Hypothetical protein.
FT NON_TER 1
SQ SEQUENCE 299 AA; 34032 MW; 6B8DB60E6A88239A CRC64;

Query Match 59.3%; Score 51; DB 4; Length 299;
Best Local Similarity 85.7%; Pred. No. 12;
Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 PMMPMPRR 11
Db 37 PMMPMSR 43

RESULT 10
P96143 PRELIMINARY; PRT; 504 AA.
ID P96143
AC P96143
DT 01-MAY-1997 (TRENBLREL. 03, Created).
DT 01-MAY-1997 (TRENBLREL. 03, Last sequence update).
DT 01-JUN-2001 (TRENBLREL. 17, Last annotation update).
DE PEPTIDE HYDROLASE.
GN TPEL.
OS Thermophilum vulgare.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Thermophilum.
NCBI_TaxID=2026;
[1]
RN SEQUENCE OF 1-431 FROM N.A.
RP STRAIN=94-2A.
RC Hofmeister J.W.;
RL Thesis (1995), Molecular Genetics,
Institut fuer Pflanzen-genetik und Kulturpflanzenforschung.
[2]
RN SEQUENCE FROM N.A.
RP STRAIN=94-2A.
RC Hofmeister J.W.;
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; Z88214; CAB0567.1; -.
DR HSSP; P00800; 1HYT.
DR Interpro; IPR001570; Peptidase_M4.
DR Pfam; PF01447; Peptidase_M4; 1.
DR Hydrolase.
KW SEQUENCE 504 AA; 56653 MW; 5A7BC05C5AD1315 CRC64;
SQ
Query Match 59.3%; Score 51; DB 2; Length 504;
Best Local Similarity 60.0%; Pred. No. 20;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
Qy 1 ILRPMWPMR 10
Db 71 LVKMTWPMR 80
RESULT 11
Q18753 PRELIMINARY; PRT; 111 AA.
ID Q18753
AC Q18753
DT 01-NOV-1996 (TRENBLREL. 01, Created).
DT 01-NOV-1996 (TRENBLREL. 01, Last sequence update).
DT 01-NOV-1998 (TRENBLREL. 08, Last annotation update).
DE GLYCINE-RICH.
GN C50P7.8
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
OC Rhabditidae; Peleodermidae; Caenorhabditis.
NCBI_TaxID=6239;
[1]
RN SEQUENCE FROM N.A.
RP MEDLINE=94150718; PubMed=7906398;
RA Wilson R., Almscough R., Anderson K., Baynes C., Berts M.,
Bontfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
Craxton M., Dear S., Du Z., Durbin R., Favello A., Fulton L.,
Gardner A., Green P., Hawkins T., Hillier L., Jier M., Johnston L.,
Jones M., Kershaw J., Kirsten J., Laister N., Latreille P.,
Lightning J., Lloyd C., Murray A., Mortimore B., O'Callaghan M.,
Parsons J., Percy C., Rifkin L., Roopra A., Saunders D., Shownkeen R.,
Smaiden N., Smith A., Sonhammer E., Staden R., Sulston J.,
Thierry-Mieg J., Thomas K., Vaubin M., Vaughan K., Waterston R.,
Watson A., Weinstock L., Wilkinson-Sproat J., Wohlman P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans";
RL Nature 368:32-38(1994).
[2]
RN SEQUENCE FROM N.A.
RP Johnson D., Stelleyes L.;
RL Submitted (DEC-1995) to the EMBL/GenBank/DBJ databases.

RN [3]
RP SEQUENCE FROM N.A.
RA Waterston R.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; U41557; AAA83303.1; -.
SQ SEQUENCE 111 AA; 10139 MW; 6E729A2E0F9762B9 CRC64;
Query Match 58.1%; Score 50; DB 5; Length 111;
Best Local Similarity 54.5%; Pred. No. 7;
Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
Qy 1 ILRPMWPMR 11
Db 12 VWRPMWPMGR 22
RESULT 12
Q9CZAI PRELIMINARY; PRT; 141 AA.
ID Q9CZAI
AC Q9CZAI
DT 01-JUN-2001 (TRENBLREL. 17, Created).
DT 01-JUN-2001 (TRENBLREL. 17, Last sequence update).
DT 01-JUN-2001 (TRENBLREL. 17, Last annotation update).
DE 2810031J10RIK PROTEIN.
GN 2810031J10RIK.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
[1]
RN SEQUENCE FROM N.A.
RP STRAIN=C57BL/6J; TISSUE=EMBRYO;
RC MEDLINE=21085660; PubMed=11217851;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yananaka I.,
Saito T., Okazaki Y., Gojohori T., Bono H., Kasukawa T., Saito R.,
Kadota K., Matsuda H.A., Ashburner M., Batilov S., Casavant T.,
Fleischmann W., Gaasterland T., Gissi C., King B., Kochia H.,
Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
Gustlich S., Hill D., Hofmann M., Hume D.A., Kamita M., Lee N.H.,
Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
Nordone P., Ring B., Ringwald C., Rodriguez I., Sakamoto N.,
Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.F.,
Suzuki H., Toyooka K., Wang K.H., Weitz C., Whitaker C., Wilming L.,
Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohlsuki S.,
Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection";
RL Nature 409:685-690(2001).
DR EMBL; AK012846; BAB28508.1; -.
DR MGI; MGI:1919917; 2810031J10RIK.
DR Interpro; IPR003309; SCAN.
DR Pfam; PF02023; SCAN; 1.
DR SMART; SM00431; LER; 1.
DR PROSITE; PS00804; SCAN_BOX; 1.
SQ SEQUENCE 141 AA; 15993 MW; 865C6B735BF8203D CRC64;
Query Match 58.1%; Score 50; DB 11; Length 141;
Best Local Similarity 66.7%; Pred. No. 8.7;
Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
Qy 1 ILRPMWPMR 9
Db 105 VSRPMWPMR 113
RESULT 13

```

Q9AUN3
ID Q9AUN3 PRELIMINARY; PRT: 327 AA.
AC Q9AUN3:
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE HYPOTHETICAL PROTEIN.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzaceae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RA Spiegel L.A., King L., Kirchoff K.A., de la Bastide M., Preston R.R.,
RA Nascimento L.U., Vil M.D., Baker J.P., Miller B., Cunnin D.M.,
RA Kult K.H., Rodriguez S., Santos L., Zutavern T., Ballja V.S.,
RA Shad R.S., Bahret A., Bal H.P., O'Shaughnessy A., Dedhia N.N.,
RA McCombie W.R.;
RT "Genomic Sequence For Oryza sativa, Nipponbare Strain, Chromosome X,
RT Clone OSJNBa0058E19, Complete Sequence.";
RL Submitted (MAR-2001) to the EMBL/Genbank/DBJ databases.
DR EMBL; AC083945; AAK13143.1;
SQ SEQUENCE 327 AA; 36672 MW; 5CCA9080664BD0CA CRC64;

Query Match 58.1%; Score 50; DB 10; Length 327;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 WMPWRR 11
| | | | |
DB 119 WMPWRR 124

RESULT 14
Q9DUC9 PRELIMINARY; PRT: 735 AA.
AC Q9DUC9:
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-MAR-2001 (TREMBLrel. 16, Last annotation update)
DE ORF1.
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=PT-TTV6;
RA Okamoto H.;
RL Submitted (Apr-2000) to the EMBL/Genbank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA STRAIN=PT-TTV6;
RX PubMed=11080484;
RA Okamoto H., Nishizawa T., Tawara A., Peng Y., Takahashi M.,
RA Kishimoto J., Tanaka T., Miyakawa Y., Mayumi M.;
RT "Species-specific TT viruses in humans and nonhuman primates and their
RT phylogenetic relatedness.";
RL Virology 277:368-378(2000).
DR EMBL; AB041957; BAB19308.1;
SQ SEQUENCE 735 AA; 86132 MW; 9ED818D6BE6FA5D3 CRC64;

Query Match 58.1%; Score 50; DB 12; Length 735;
Best Local Similarity 46.7%; Pred. No. 37;
Matches 7; Conservative 2; Mismatches 2; Indels 4; Gaps 1;

QY 2 LRPWPM-----WPMRRK 12
: | | | |
DB 1 MAMPWRRRRRRRRRR 15

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RESULT 15
Q9DT80 PRELIMINARY; PRT: 49 AA.
AC Q9DT80:
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DE 01-MAR-2001 (TREMBLrel. 16, Last annotation update)
DE ORF1 (FRAGMENT).
OS TT virus.
OC Viruses; ssDNA viruses; unclassified ssDNA viruses.
OX NCBI_TaxID=68887;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=TYM9;
RX MEDLINE=20568739; PubMed=1118348;
RA Okamoto H., Nishizawa T., Tawara A., Takahashi M., Kishimoto J.,
RA Sai T., Sugai Y.;
RT "TT virus mRNAs detected in the bone marrow cells from an infected
RT individual.";
RL Biochem. Biophys. Res. Commun. 279:700-707(2000).
DR EMBL; AB050449; BAB19930.1;
FT NON_TER 49
SQ SEQUENCE 49 AA; 7225 MW; 1DA6F81AB69AA43 CRC64;

Query Match 57.0%; Score 49; DB 12; Length 49;
Best Local Similarity 36.8%; Pred. No. 4.6;
Matches 7; Conservative 2; Mismatches 2; Indels 8; Gaps 1;

QY 2 LRPWPM-----WPMRRK 12
: | | | |
DB 1 MAMPWRRRRRRRRRR 19

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Search completed: January 4, 2002, 08:47:26
Job time: 414 sec

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 4, 2002, 08:47:48 ; Search time 18.1 Seconds

(without alignments)
24.308 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRMPMPWRRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 100059 segs, 36664827 residues

Total number of hits satisfying chosen parameters: 100059

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_39.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	70	81.4	144	1	INDC_BOVIN	P33046 bos taurus
2	53	61.6	1173	1	VGL2_CVH22	P15423 human coron
3	49	57.0	492	1	ADRO_BOVIN	P08165 bos taurus
4	47	54.7	253	1	Y945_MYCTU	P71564 mycobacteri
5	47	54.7	715	1	YD55_MYCTU	Q11025 mycobacteri
6	45.5	52.9	505	1	TRPE_PSRSS	P21689 pseudomonas
7	45	52.3	196	1	YA05_SCHPO	Q09677 schizosacch
8	45	52.3	1108	1	CN38_RAT	O63083 rattus norv
9	44	51.2	361	1	FUT3_HUMAN	P12121 homo sapien
10	44	51.2	372	1	FUT3_PANTR	Q19058 pan troglod
11	44	51.2	535	1	MM6_MYCTU	Q10773 mycobacteri
12	44	51.2	535	1	YD6_SCHPO	Q13912 schizosacch
13	44	51.2	967	1	MM4_MYCTU	O53738 mycobacteri
14	44	51.2	968	1	MM2_MYCTU	Q11171 mycobacteri
15	44	51.2	984	1	SLX3_MOUSE	Q04891 mus musculu
16	44	51.2	1154	1	VGL2_IBVD2	P12722 avian infec
17	44	51.2	1162	1	VGL2_IBVK	P11223 avian infec
18	44	51.2	1162	1	VGL2_IBVK	P12650 avian infec
19	44	51.2	1162	1	VGL2_IBVM	P12651 avian infec
20	44	51.2	1163	1	VGL2_IBV6	P05135 avian infec
21	43.5	50.6	276	1	KCE1_RHOP	O83005 rhodospseud
22	43.5	50.6	2436	1	ABC2_HUMAN	Q09267 homo sapien
23	43	50.0	51	1	LHB2_ECTHA	P01005 ectothiorho
24	43	50.0	711	1	MM1A_STRCO	O53902 streptomyc
25	43	50.0	958	1	MM1A_MYCTU	P95211 mycobacteri
26	43	50.0	1112	1	CN3B_HUMAN	Q13370 homo sapien
27	43	50.0	1225	1	VGL2_CVPR8	P27653 porcine res
28	43	50.0	1225	1	VGL2_CVPRM	P24413 porcine res
29	43	50.0	1235	1	VGL2_CVMAH	P11225 murine coro
30	43	50.0	1324	1	VGL2_CVMA5	P11224 murine coro
31	43	50.0	1353	1	VGL2_CVHOC	P36334 human coron
32	43	50.0	1363	1	VGL2_CVPR	P25190 bovine coro
33	43	50.0	1363	1	VGL2_CVBL9	P25191 bovine coro

34	43	50.0	1363	1	VGL2_CVBLX	P25192 bovine coro
35	43	50.0	1363	1	VGL2_CVBM	P15777 bovine coro
36	43	50.0	1363	1	VGL2_CVBO	P25193 bovine coro
37	43	50.0	1363	1	VGL2_CVBP	P25194 bovine coro
38	43	50.0	1376	1	VGL2_CVMA	P22432 murine coro
39	43	50.0	1376	1	VGL2_CVMC	Q02385 murine coro
40	43	50.0	1447	1	VGL2_CVPR	Q02167 porcine tra
41	43	50.0	1447	1	VGL2_CVPRU	P07946 porcine tra
42	43	50.0	1447	1	VGL2_CVPRT	Q01977 porcine tra
43	43	50.0	1449	1	VGL2_CVPRS	P18450 porcine tra
44	43	50.0	1449	1	VGL2_CVPR	P33470 porcine tra
45	43	50.0	1451	1	VGL2_CVCAI	P36300 canine ente

ALIGNMENTS

RESULT 1	INDC_BOVIN	STANDARD	PRT	144 AA.
AC	P33046:			
DT	01-OCT-1993 (Rel. 27, Created)			
DT	01-OCT-1993 (Rel. 27, Last sequence update)			
DT	01-NOV-1997 (Rel. 35, Last annotation update)			
DE	INDOLICIDIN PRECURSOR.			
OS	Bos taurus (Bovine).			
OC	Eukaryota: Metazoa: Chordata: Vertebrata: Euteleostomi:			
OC	Mammalia: Eutheria: Cetartiodactyla: Ruminantia: Pecora: Bovidea:			
OC	Bovidae: Bovinae: Bos.			
OX	NCBI_TaxID:9913;			
RP	(1)			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Bone marrow;			
RX	MEDLINE=92392368; PubMed=1520337;			
RA	del Sal G., Storici P., Schneider C., Romeo D., Zanetti M.;			
RT	"cDNA cloning of the neutrophil bactericidal peptide indolicidin.";			
RL	Biochem. Biophys. Res. Commun. 187:467-472(1992).			
RN	[2]			
RP	SEQUENCE OF 131-143.			
RC	TISSUE=Neutrophils;			
RX	MEDLINE=92165771; PubMed=1537821;			
RA	Seisted M.E., Novotny M.J., Morris W.L., Tang Y.-O., Smith W.;			
RT	"Indolicidin, a novel bactericidal tridecapeptide amide from			
RL	neutrophils.";			
RP	J. Biol. Chem. 267:4292-4295(1992).			
CC	-1- FUNCTION: POTENT MICROBICIDAL ACTIVITY, ACTIVE AGAINST			
CC	STAPHYLOCOCCUS AUREUS AND ESCHERICHIA COLI.			
CC	-1- TISSUE SPECIFICITY: LARGE GRANULES OF NEUTROPHILS.			
CC	-1- PWM: ELASTASE MIGHT BE RESPONSIBLE FOR ITS MATURATION.			
CC	-1- SIMILARITY: BELONGS TO THE CATHELICIDIN FAMILY.			
CC	-----			
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CC	between the Swiss Institute of Bioinformatics and the EMBL outstation -			
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CC	entities requires a license agreement (see http://www.isb-sib.ch/announce/			
CC	or send an email to license@sib-sib.ch).			
CC	-----			
DR	EMBL: X67340; CAA47755.1; -			
DR	PIR: JCI222; JCI222.			
DR	PIR: A42387; A42387.			
DR	InterPro: IPR001894; Cathelicidin.			
DR	Pfam: PF00666; Cathelicidins; 1.			
DR	ProDom: PD001838; Cathelicidin; 1.			
DR	PROSITE: PS00946; CATHELICIDINS_1; 1.			
DR	PROSITE: PS00947; CATHELICIDINS_2; 1.			
KW	Antibiotic; Amidation; Signal.			
FT	SIGNAL	1	29	POTENTIAL.
FT	PROPEP	30	130	
FT	PEPTIDE	131	143	INDOLICIDIN, CARBOXYLIC ACID (BY
FT	MOD_RES	30	30	PYRROLIDONE

FT	DISULFID	85	96		SIMILARITY.
FT	DISULFID	107	124		BY SIMILARITY.
FT	MOD_RES	143	143		AMIDATION (G-144 PROVIDE AMIDE GROUP).
SQ	SEQUENCE	144 AA;	16479 MW;	E3B1CBEB5C0911 CRC64;	
Query Match		Best Local Similarity	81.4%;	Score 70; DB 1;	Length 144;
Matches	8; Conservative		Pred. No. 0.0039;	Mismatches 0;	Indels 0; Gaps 0;
OY	3 RWPMPWRR	11			
Dd	135 KWPMPWRR	143			
RESULT	2				
ID	VGL2_CVH22	STANDARD;	PRT:	1173 AA.	
AC	P15423;				
DT	01-APR-1990 (Rel. 14, Created)				
DT	01-APR-1990 (Rel. 14, Last sequence update)				
DT	15-JUN-1999 (Rel. 38, Last annotation update)				
DE	E2 GLYCOPROTEIN PRECURSOR (SPIKE GLYCOPROTEIN) (PEPLOMER PROTEIN).				
GN	S.				
OS	Human coronavirus (strain 229E).				
OC	Viruses; ssRNA positive-strand viruses, no DNA stage; Nidovirales;				
OX	Coronaviridae; Coronavirus.				
OX	NCBI_TaxID=11137;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=90264837; PubMed=2245367;				
RA	Raabe T., Schelle-Prinz B., Siddell S.G.;				
RT	"Nucleotide sequence of the gene encoding the spike glycoprotein of human coronavirus HCoV 229E."				
RL	J. Gen. Virol. 71:1065-1073(1990).				
CC	-1- FUNCTION: THE PELOMER PROTEIN MEDIATES THE BINDING OF VIRIONS TO THE HOST CELL RECEPTOR AND IS INVOLVED IN MEMBRANE FUSION				
CC	AND IN STINCTUUM FORMATION.				
CC	-1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.				
CC	-----				
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CC	-----				
DR	EMBL: X16816; CAA34723.1;				
DR	PIR: A34766; VGIHHC.				
DR	InterPro: IPRO02551; Corona_S1.				
DR	InterPro: IPRO02552; Corona_S2.				
DR	Pfam: PF01600; Corona_S1; 1.				
DR	Pfam: PF01601; Corona_S2; 1.				
KW	Glycoprotein; Envelope protein; Transmembrane; Signal.				
FT	SIGNAL	1	15		
FT	CHAIN	16	1173		E2 GLYCOPROTEIN.
FT	DOMAIN	16	1115		EXTRACELLULAR (POTENTIAL).
FT	TRANSMEM	1116	1135		POTENTIAL.
FT	DOMAIN	1136	1173		CYTOSOLASMIC (POTENTIAL).
FT	DOMAIN	1136	1157		CYS-RICH.
FT	CARBOHYD	23	23		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	62	62		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	98	98		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	147	147		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	171	171		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	176	176		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	220	220		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	243	243		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	326	326		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	333	333		N-LINKED (GLCNAG . . .) (POTENTIAL).
FT	CARBOHYD	440	440		N-LINKED (GLCNAG . . .) (POTENTIAL).

Query Match	Best Local Similarity	Score	DB 1:	Length	Indels	Gaps		
Matches	5;	Conservative	2;	Mismatches	1;	Indels	0;	Gaps
QY	2	LRPMPWPW 9	:::					
Db	1112	IKPMPWVW 1119						
RESULT	3							
ADRO_BOVIN	STANDARD;	PRT;	492	AA.				
AC	P08165;							
DT	01-AUG-1988 (Rel. 08, Created)							
DT	15-JUL-1998 (Rel. 36, Last sequence update)							
DT	20-AUG-2001 (Rel. 40, Last annotation update)							
DE	NADPH:ADRENODOXIN OXIDOREDUCTASE, MITOCHONDRIAL, PRECURSOR							
DE	(EC 1.18.1.2) (ADRENODOXIN REDUCTASE) (AR) (FERREDOXIN-NADP(+) REDUCTASE).							
GN	FDXR OR ADXR.							
OS	Bos taurus (Bovine).							
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;							
OC	Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;							
OC	Bovidae; Bovinae; Bos.							
OX	NCBI_TaxID=9913;							
RN	[1]							
RP	SEQUENCE FROM N.A., AND ALTERNATIVE SPLICING.							
RX	MEDLINE=94177140; PubMed=8130767;							
RA	Takata Y., Sagara Y., Kono A., Sekimizu K., Horiiuchi T.;							
RT	"Gene structure of bovine adrenodoxin reductase.";							
RL	Biol. Pharm. Bull. 16:1200-1206(1993).							
RN	[2]							
RP	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.							
RX	MEDLINE=88198050; PubMed=448086;							
RA	Sagara Y., Takata Y., Miyata T., Hara T., Horiiuchi T.;							
RT	"Cloning and sequence analysis of adrenodoxin reductase cDNA from							
RL	bovine adrenal cortex.";							
RN	J. Biochem. 102:1333-1336(1987).							
RN	[3]							
RP	SEQUENCE FROM N.A.							
RX	MEDLINE=87270696; PubMed=3038094;							
RA	Nonaka Y., Murakami H., Yabusaki Y., Kuramitsu S., Kagamiyama H.,							
RT	Yanano T., Okamoto M.;							
RL	"Molecular cloning and sequence analysis of full-length cDNA for							
RN	of adrenodoxin oxidoreductase from bovine adrenal cortex.";							
RL	Biochem. Biophys. Res. Commun. 145:1239-1247(1987).							
RN	[4]							
RP	SEQUENCE FROM N.A.							
RC	TISSUE=Adrenal cortex;							
RX	MEDLINE=89170752; PubMed=2924777;							
RA	Hankoglu I., Gutfinger T.;							

RT "cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites in oxidoreductases.";
 RT Eur. J. Biochem. 180:479-484(1989).
 RN [5]
 RP SEQUENCE OF N-TERMINUS, AND PARTIAL SEQUENCE.
 RC TISSUE-Adrenal cortex;
 RX MEDLINE=88082777; PubMed=3691502;
 RA Hanukoglu I., Gutfinger T., Hanlu M., Shively J.E.;
 RT "Isolation of a cDNA for adrenodoxin reductase (ferredoxin-NADP+ reductase). Implications for mitochondrial cytochrome P-450 systems.";
 RL Eur. J. Biochem. 169:449-455(1987).
 RN [6]
 RP X-RAY CRYSTALLOGRAPHY (1.7 ANGSTROMS) OF 33-492.
 RC TISSUE-Adrenal gland;
 RX MEDLINE=99299392; PubMed=10369776;
 RA Ziegler G.A., Vonnheim C., Hanukoglu I., Schulz G.E.;
 RT "The structure of adrenodoxin reductase of mitochondrial P450 systems: electron transfer for steroid biosynthesis.";
 RL J. Mol. Biol. 289:981-990(1999).
 CC -1- FUNCTION: SERVES AS THE FIRST ELECTRON TRANSFER PROTEIN IN ALL THE MITOCHONDRIAL P450 SYSTEMS, INCLUDING CHOLESTEROL SIDE CHAIN CLEAVAGE IN ALL STEROIDGENIC TISSUES, STEROID 11-BETA HYDROXYLATION IN THE ADRENAL CORTEX, 25-OH-VITAMIN D3-24 HYDROXYLATION IN THE KIDNEY, AND STEROL C-27 HYDROXYLATION IN THE LIVER.
 CC -1- CATALYTIC ACTIVITY: REDUCED ADRENODOXIN + NADP(+) = OXIDIZED ADRENODOXIN + NADPH.
 CC -1- COFACTOR: FAD.
 CC -1- PATHWAY: CHOLESTEROL SIDE-CHAIN-CLEAVAGE SYSTEM.
 CC -1- SUBUNIT: MONOMER.
 CC -1- SUBCELLULAR LOCATION: MITOCHONDRIAL MATRIX.
 CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS. A SHORT FORM (SHOWN HERE) AND A LONG FORM. ARE PRODUCED BY ALTERNATIVE SPLICING. THE LONG FORM REPRESENTS 10-20% OF ALL ADRENODOXIN REDUCTASE MRNA. AND SEEMS TO BE INACTIVE.
 CC CC
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 CC -----
 DR EMBL: M17029; AAA0362.1; -
 DR EMBL: D00211; BA00150.1; -
 DR EMBL: X13736; CA032002.1; -
 DR PIR: A29604; A29604.
 DR PIR: JS0390; JS0390.
 DR PIR: S03558; S03558.
 DR PIR: J70751; J70751.
 DR PDB: 1CJC; 12-APR-99.
 DR PDB: 1E1L; 02-JUN-00.
 DR InterPro: IPR000759; Adnrdx_redctase.
 DR PRINTS: PR00419; ADXRDPASE.
 KM Electron transport: Oxidoreductase; Flavoprotein; NADP: FAD; Mitochondrion; Transit peptide; Alternative splicing; 3D-structure.
 FT TRANSIT 1 32 MITOCHONDRION.
 FT CHAIN 33 492 NADPH:ADRENODOXIN OXIDOREDUCTASE.
 FT VARSPLIC 204 204 E -> EVLLICQ (IN LONG ISOFORM).
 FT CONFLICT 77 77 G -> R (IN REF. 3).
 FT CONFLICT 81 94 FGVARDHPEVKNI -> VWLALTPPRSMIL (IN REF. 3).
 FT CONFLICT 124 128 QDAYH -> RYRRLT (IN REF. 3).
 FT CONFLICT 268 268 K -> R (IN REF. 3).
 FT CONFLICT 317 318 PS -> RL (IN REF. 3).
 FT CONFLICT 323 333 RAAGIRLAVTR -> ARSASMSPE (IN REF. 3).
 FT CONFLICT 341 352 TRAVPTGVEDL -> HFGSAHWGCGP (IN REF. 3).
 SQ SEQUENCE 492 AA; 54338 MW; E68F6F5D18F53131 CRC64;

Query Match
 Best Local Similarity

57.0%; Score 49; DB 1; Length 492;
 83.3%; Pred. No. 6.6;

Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 4 WPMWPM 9
 Db 6 WRWPMW 11

RESULT 4

Y945_MYCTU STANDARD; PRT; 253 AA.
 AC P71564;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE OXIDOREDUCTASE RV0945 (EC 1.-.-.-).
 GN RV0945 OR MT0971 OR MYC110D7.29C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OX NCBI_TaxID=1773;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D., Gordon S.V., Eigmeier K., Gas S., Barry C.E. III, Tekala F., Badcock K., Basham D., Brown D., Chillingworth T., Connor R., Davies R., Devlin K., Felwell T., Gentles S., Hamlin N., Holroyd S., Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy J., Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J., Rutter S., Seeger K., Skelton S., Squares R., Sultun J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.";
 RT Nature 393:537-544(1998).
 RL [2]
 RN SEQUENCE FROM N.A.
 RP STRAIN=CDC 1551 / Oshkosh;
 RC Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O., Peterson S., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E., Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L., Delcher A., Utterback T., Weidman J., Khouri H., Gill J., Mikula A., Bishal W.;
 RA "Whole genome comparison of Mycobacterium tuberculosis clinical and laboratory strains.";
 RT Submitted (Apr-2001) to the EMBL/Genbank/DBJ databases.
 RL -1- SIMILARITY: BELONGS TO THE SHORT-CHAIN DEHYDROGENASES/REDUCTASES (SDR) FAMILY.
 CC CC

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 CC -----

DR EMBL: Z79700; CAB02005.1; -
 DR EMBL: AE006982; AAK45219.1; -
 DR TIGR: MT0971; -
 DR Tuberculist: RV0945; -
 DR InterPro: IPR002198; ADH_SHORT.
 DR Pfam: PF00106; adh_short.1.
 DR PROSITE: PS00061; ADH_SHORT; 1.
 DR Hypothetical protein; Oxidoreductase; Complete proteome.
 FT ACT_SITE 159 159 BY SIMILARITY.
 SQ SEQUENCE 253 AA; 27138 MW; BAD937208842DA12 CRC64;

Query Match
 Best Local Similarity
 Matches 5; Conservative

54.7%; Score 47; DB 1; Length 253;
 100.0%; Pred. No. 6.6;
 0; Mismatches 0; Indels 0; Gaps 0;

RP SEQUENCE FROM N.A.
 RC STRAIN-972;
 RA Connor R., Churcher C.M., Barrell B.G., Rajandream M.A., Walsh S.V.;
 RL Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases.
 CC -1 SIMILARITY: STRONG, TO BACTERIAL MODULATOR OF DRUG ACTIVITY B
 (MDAB).
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 CC -----
 CC EMBL: Z49811; CAAB9955.1;
 DR InterPro: IPR003680; NADHdh_2;
 DR Pfam: PF02525; NADHdh_2; 1.
 KM Hypothetical protein.
 SQ SEQUENCE 196 AA; 22104 MW; 436764DA9E26074C CRC64;
 Query Match 52.38; Score 45; DB 1; Length 196;
 Best Local Similarity 50.0%; Pred. No. 9.5;
 Matches 8; Conservative 2; Mismatches 2; Indels 4; Gaps 2;
 QY 1 ILRMP-WW---PWRRK 12
 Db 63 IYQWPGWMMCTPWRLK 78
 RESULT 8
 CN3B_RAT STANDARD; PRT: 1108 AA.
 ID Q63085;
 AC 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE GMP-INHIBITED 3', 5'-CYCLIC PHOSPHODIESTERASE B (EC 3.1.4.17) (CYCLIC
 DE GMP INHIBITED PHOSPHODIESTERASE B) (CGI-PDE B) (CGIPDEB).
 GN PDE3B.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxId=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-SPRAGUE-DAWLEY; TISSUE=adipose tissue;
 RX MEDLINE=93366761; PubMed=8395509;
 RA Taira M., Hockman S.C., Calvo J.C., Taira M., Belfrage P.,
 RA Mangiatello V.C.;
 RT "Molecular cloning of the rat adipocyte hormone-sensitive cyclic GMP-
 RT inhibited cyclic nucleotide phosphodiesterase.";
 RL J. Biol. Chem. 268:18573-18579(1993).
 CC -1 FUNCTION: MAY PLAY A ROLE IN FAT METABOLISM.
 CC -1 CATALYTIC ACTIVITY: GUANOSINE 3', 5'-CYCLIC PHOSPHATE + H(2)O =
 CC GUANOSINE 5'-PHOSPHATE.
 CC -1 ENZYME REGULATION: INHIBITED BY GMP.
 CC -1 SUBCELLULAR LOCATION: MEMBRANE-BOUND (POTENTIAL).
 CC -1 TISSUE SPECIFICITY: ABUNDANT IN ADIPOSE TISSUES.
 CC -1 SIMILARITY: BELONGS TO THE CYCLIC NUCLEOTIDE PHOSPHODIESTERASE
 CC FAMILY.
 CC -----
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 CC -----
 CC EMBL: Z22867; CAAB0489.1;
 DR InterPro: IPR003607; HDC.

DR InterPro: IPR002073; PDEase.
 DR Pfam: PF00233; PDEase; 1.
 DR SMART: SM00471; HDC; 1.
 DR PROSITE: PS00126; PDEASE_I; 1.
 KM Hydrolase; GMP; Membrane.
 FT DOMAIN 16 22 POLY-PRO.
 FT DOMAIN 99 102 POLY-ALA.
 FT DOMAIN 175 179 POLY-ALA.
 FT DOMAIN 1007 1021 POLY-ASP.
 FT DOMAIN 1068 1071 POLY-GLU.
 FT DOMAIN 1101 1104 POLY-GLU.
 SQ SEQUENCE 1108 AA; 123105 MW; C9B5078C7D3ADD6D CRC64;
 Query Match 52.38; Score 45; DB 1; Length 1108;
 Best Local Similarity 62.5%; Pred. No. 46;
 Matches 5; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
 QY 4 WPMWRR 11
 Db 164 WQWWSWLR 171
 RESULT 9
 FUT3_HUMAN STANDARD; PRT: 361 AA.
 ID P21217; Q09448; Q09449.
 AC 01-MAY-1991 (Rel. 18, Created)
 DT 01-MAY-1991 (Rel. 18, Last sequence update)
 DT 30-MAY-1991 (Rel. 39, Last annotation update)
 DE GALACTOSIDE 3(4)-L-FUCOSYLTRANSFERASE (EC 2.4.1.65) (BLOOD GROUP LEWIS
 DE ALPHA-4-FUCOSYLTRANSFERASE) (LEWIS FT) (FUCOSYLTRANSFERASE 3) (FUCT-
 DE III).
 GN FUT3 OR LE.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_TaxId=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91032981; PubMed=1977660;
 RA Kukowska-latallo J.F., Larsen R.D., Nair R.P., Lowe J.B.;
 RT "A cloned human cDNA determines expression of a mouse stage-specific
 RT embryonic antigen and the Lewis blood group
 RT alpha(1,3/1,4)fucosyltransferase.";
 RL Genes Dev. 4:1288-1303(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;
 RX MEDLINE=95378269; PubMed=7650030;
 RA Cameron H.S., Szecepaniak D., Weston M.;
 RT "Expression of human chromosome 19p alpha(1,3)-fucosyltransferase
 RT genes in normal tissues. Alternative splicing, polyadenylation, and
 RT isoforms.";
 RL J. Biol. Chem. 270:20112-20122(1995).
 RN [3]
 RP VARIANT LE(-) MET-105.
 RX MEDLINE=94059067; PubMed=8240322;
 RA Elmgren A., Rydberg L., Larson G.;
 RT "Genocytic heterogeneity among Lewis negative individuals.";
 RL Biochem. Biophys. Res. Commun. 196:515-520(1993).
 RN [4]
 RP VARIANTS LE(-) ARG-20; SER-170 AND ALA-336.
 RX MEDLINE=94059082; PubMed=8240337;
 RA Nishihara S., Iazawa S., Iwasaki H., Nakazato M., Kudo T., Ando T.,
 RA Nishimatsu H.;
 RT "Alpha (1,3/1,4)fucosyltransferase (FUCT-III) gene is inactivated by
 RT a single amino acid substitution in Lewis histo-blood type negative
 RT individuals.";
 RL Biochem. Biophys. Res. Commun. 196:624-631(1993).
 RN [5]
 RP VARIANTS LE(-) ARG-20 AND SER-170.
 RX MEDLINE=94033579; PubMed=8219240;

RA Koda Y., Kimura H., Mekada E.:
 RT "Analysis of Lewis fucosyltransferase genes from the human gastric
 RT mucosa of Lewis-positive and -negative individuals.";
 RL Blood 82:2915-2919(1993).
 RN [6]
 RP VARIANTS LE(-) ARG-20 AND LYS-356.
 RX MEDLINE=94342259; PubMed=8063716;
 RA Mollitone R., Reguigne I., Kelly R.J., Fletcher A., Watt J.,
 RA Chatfield S., Aziz A., Cameron H.S., Weston B.W., Lowe J.B., Oriol R.:
 RT "Molecular basis for Lewis alpha(1,3/1,4)-fucosyltransferase gene
 RT deficiency (FUT3) found in Lewis-negative Indonesian pedigrees.";
 RL J. Biol. Chem. 269:20987-20994(1994).
 RN [7]
 RP VARIANT LE(-) LYS-356.
 RX MEDLINE=95050753; PubMed=7961897;
 RA Nishihara S., Narimatsu H., Iwasaki H., Yazawa S., Akamatsu S.,
 RA Ando T., Seno T., Narimatsu I.:
 RT "Molecular genetic analysis of the human Lewis histo-blood group
 RT system.";
 RL J. Biol. Chem. 269:29271-29278(1994).
 RN [8]
 RP VARIANTS LE(-) ARG-20; ARG-68; MET-105 AND LYS-356.
 RX MEDLINE=96243526; PubMed=8801770;
 RA Elmgren A., Boerjeson C., Svensson L., Rydberg L., Larson G.:
 RT "DNA sequencing and screening for point mutations in the human Lewis
 RT 'FUT3' gene enables molecular genotyping of the human Lewis blood
 RT group system.";
 RL Vox Sang. 70:97-103(1996).
 RN [9]
 RP VARIANTS LE(-) ARG-68 AND MET-105.
 RX MEDLINE=97413801; PubMed=9268337;
 RA Elmgren A., Mollitone R., Costache M., Boerjeson C., Oriol R.,
 RA Harrington J., Larson G.:
 RT "Significance of individual point mutations, T202C and C314T, in the
 RT human Lewis 'FUT3' gene for expression of Lewis antigens by the human
 RT alpha1,3/1,4-fucosyltransferase, Fuc-TIII.";
 RL J. Biol. Chem. 272:21994-21998(1997).
 RN [10]
 RP VARIANTS LE(+) K-102; A-124, AND VARIANTS LE(-) N-162; R-223; M-270.
 RX MEDLINE=98366989; PubMed=9703429;
 RA Pang H., Liu Y., Koda Y., Soejima M., Jia J., Schlaphoff T.,
 RA du Toit E.D., Kimura H.:
 RT "Five novel missense mutations of the Lewis gene 'FUT3' in African
 RT 'Xhosa' and Caucasian populations in South Africa.";
 RL Hum. Genet. 102:675-680(1998).
 CC -I- FUNCTION: MAY CATALYZE ALPHA-1,3 AND ALPHA-1,4 GLYCOSIDIC LINKAGES
 CC INVOLVED IN THE EXPRESSION OF VIM-2, LEWIS A, LEWIS B, STAYL
 CC LEWIS X AND LEWIS X/SSA-1 ANTIGENS. MAY BE INVOLVED IN BLOOD
 CC GROUP LEWIS DETERMINATION; LEWIS-POSITIVE (LE(+)) INDIVIDUALS
 CC HAVE AN ACTIVE ENZYME WHILE LEWIS-NEGATIVE (LE(-)) INDIVIDUALS
 CC HAVE AN INACTIVE ENZYME.
 CC -I- CATALYTIC ACTIVITY: GDP-L-FUCOSE + 1,3-BETA-D-GALACTOSYL-
 CC N-ACETYL-D-GLUCOSAMINYL-R = GDP + 1,3-BETA-D-GALACTOSYL-
 CC (ALPHA-1,4-L-FUCOSYL)-N-ACETYL-D-GLUCOSAMINYL-R.
 CC -I- PATHWAY: GLYCOSYLATION.
 CC -I- SUBCELLULAR LOCATION: TYPE II MEMBRANE PROTEIN. MEMBRANE-BOUND
 CC FORM IN TRANS CISTERNAE OF GOLGI.
 CC -I- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN STOMACH, COLON, SMALL
 CC INTESTINE, LUNG AND KIDNEY AND TO A LESSER EXTENT IN SALIVARY
 CC GLAND, BLADDER, UTERUS AND LIVER.
 CC -I- MISCELLANEOUS: ALSO ACTS ON THE CORRESPONDING 1,4-GALACTOSYL
 CC DERIVATIVE, FORMING 1,3-L-FUCOSYL LINKS.
 CC -I- SIMILARITY: STRUCTURAL SIMILARITY WITH THE OTHER MAMMALIAN
 CC GLYCOSYLTRANSFERASES.
 CC -----
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 CC -----

DR EMBL: X53578; CAA37641.1; -
 DR EMBL: U27328; AAC50187.1; -
 DR EMBL: U27326; AAC50185.1; -
 DR EMBL: U27327; AAC50186.1; -
 DR EMBL: D89324; BAA13941.1; -
 DR EMBL: D89325; BAA13942.1; -
 DR PIR: A36669; A36669.
 DR MIM: 111100; -
 DR InterPro: IPR001503; Glyco_transf_10.
 DR Pfam: PF00852; Glyco_transf_10.1.
 KW Transferase; Glycosyltransferase; Glycoprotein; Transmembrane;
 KW Signal-anchor; Golgi stack; Polymorphism; Blood group antigen.
 KW Cytoplasmic (POTENTIAL).
 KW DOMAIN 1 15
 FT TRANSMEM 16 34
 FT DOMAIN 35 361
 FT CARBOHYD 154 154
 FT CARBOHYD 185 185
 FT VARIANT 20 20
 FT VARIANT 68 68
 FT VARIANT 102 102
 FT VARIANT 105 105
 FT VARIANT 124 124
 FT VARIANT 162 162
 FT VARIANT 170 170
 FT VARIANT 223 223
 FT VARIANT 270 270
 FT VARIANT 336 336
 FT VARIANT 356 356
 FT SEQUENCE 361 AA; 42117 MW; Bf4398044F19C284 CRC64;
 SQ
 Query Match 51.2%; Score 44; DB 1; Length 361;
 Best Local Similarity 85.7%; Pred. No. 23;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 5 PWWPWR 11
 DB 9 PWWPWR 15
 RESULT 10
 ID FUT3_PANTR STANDARD; PRT; 372 AA.
 AC 019058;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 30-MAY-2000 (Rel. 39, Last annotation update)
 DE GALACTOSIDE 3(4)-L-FUCOSYLTRANSFERASE (EC 2.4.1.65) (BLOOD GROUP LEWIS
 DE ALPHA-4-FUCOSYLTRANSFERASE) (LEWIS FT) (FUCOSYLTRANSFERASE 3) (FUCT-
 DE IIT) (ALPHA-3/4-FUCOSYLTRANSFERASE).
 GN FUT3.
 OS Pan troglodytes (Chimpanzee).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pan.
 OX NCBI_TaxID=9598;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=98037800; PubMed=9368041;
 RA Costache M., Apoll P.-A., Cailliau A., Elmgren A., Larson G.,
 RA Henry S., Blanche A., Iordachescu D., Oriol R., Mollitone R.:
 RT "Evolution of fucosyltransferase genes in vertebrates.";

RL J. Biol. Chem. 272:29721-29728(1997).
 CC -1- FUNCTION: MAY CATALYZE ALPHA-1,3 AND ALPHA-1,4 GLYCOSIDIC LINKAGES
 CC INVOLVED IN THE EXPRESSION OF STAIYL LEMIS X AND LEMIS X/SEEA-1
 CC ANTIGENS. IT MAY BE INVOLVED IN BLOOD GROUP LEMIS DETERMINATION
 CC (BY SIMILARITY).
 CC -1- CATALYTIC ACTIVITY: GDP-L-FUCOSE + 1,3-BETA-D-GALACTOSYL-
 CC N-ACETYL-D-GLUCOSAMINYL-R = GDP + 1,3-BETA-D-GALACTOSYL-
 CC (ALPHA-1,4-L-FUCOSYL)-N-ACETYL-D-GLUCOSAMINYL-R.
 CC -1- PATHWAY: GLYCOSYLATION.
 CC -1- SUBCELLULAR LOCATION: TYPE II MEMBRANE PROTEIN. MEMBRANE-BOUND
 CC FORM IN TRANS CISTERNAE OF GOLGI (BY SIMILARITY).
 CC -1- POLYMORPHISM: THERE ARE TWO ALLELES (A AND B). ALLELE A HAS ARG-
 CC 162 AND VAL-304. ALLELE B HAS GLY-162 AND MET-304.
 CC -1- MISCELLANEOUS: ALSO ACTS ON THE CORRESPONDING 1,4-GALACTOSYL
 CC DERIVATIVE, FORMING 1,3-L-FUCOSYL LINKS.
 CC -1- SIMILARITY: STRUCTURAL SIMILARITY WITH THE OTHER MAMMALIAN
 CC GLYCOSYLTRANSFERASES.
 CC -----
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 CC -----
 DR EMBL: Y14033; CAA74360.1; -
 DR InterPro: IPR001503; Glyco_transf_10.
 DR Pfam: PF00852; Glycoyltransferase; Glycoprotein; Transmembrane;
 KW Transferase; Glycosyltransferase; Glycoprotein; Transmembrane;
 FT Signal-anchor; Golgi stack; Polymorphism.
 FT DOMAIN 1 14 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 15 34 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
 FT (POTENTIAL).
 FT DOMAIN 35 372 LUMENAL, CATALYTIC (POTENTIAL).
 FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 196 196 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT VARIANT 162 162 R -> G (IN ALLELE B).
 FT VARIANT 304 304 V -> M (IN ALLELE B).
 FT SEQUENCE 372 AA; 43233 MW; 649CB8BCA7BD74C CRC64;

Query Match 51.2%; Score 44; DB 1; Length 372;
 Best Local Similarity 85.7%; Pred. No. 23;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 RWPMPWR 11
 I I I I I
 Db 9 RWPMPWR 15

RESULT 11
 MML6_MYCTU STANDARD; PRT; 397 AA.
 AC Q10773;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, last sequence update)
 DT 20-AUG-2001 (Rel. 40, last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML6.
 GN MML6 OR RV157 OR MT1608 OR MTCY48.08C.
 OS Mycobacterium tuberculosis.
 CC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 CC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
 CC NCI_TaxID=1773;
 RX SEQUENCE FROM N.A.
 RP STRAIN-H37RV;
 RC MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 RA Gordon S.V., Eiglmeier K., Gas S., Barry C.E. III, Tekle A.F.,
 RA Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 RA Davies R., Devlin K., Fellwell T., Gentles S., Hamlin N., Holroyd S.,
 RA Hornsby T., Jagers K., Krogh A., McLean J., Moule S., Murphy L.,

RA Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 RA Ruter S., Seeger K., Skelton S., Squares S., Squares R.,
 RA Sulten J.E., Taylor K., Whitehead S., Barrell B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-CDC 1551 / Oshkosh;
 RA Fleischmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
 RA Peterson J., Deboy R., Dodson R., Gwinn M.L., Hatt D., Hickey E.,
 RA Kolonay J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
 RA Kohnen A., Uetebach T., Weidman J., Khouri H., Gill J., Mikula A.,
 RA Bishai W.;
 RT "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RT laboratory strains."
 RL Submitted (APR-2001) to the EMBL/Genbank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML6 FAMILY.
 CC -----
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 CC -----
 DR EMBL: Z74020; CAA98334.1; -
 DR EMBL: AE007027; AAK45875.1; -
 DR TIGR: MT1608; -
 DR TubercuList; RV1557; -
 KW Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 161 181 POTENTIAL.
 FT TRANSMEM 190 210 POTENTIAL.
 FT TRANSMEM 214 234 POTENTIAL.
 FT TRANSMEM 242 262 POTENTIAL.
 FT TRANSMEM 293 313 POTENTIAL.
 FT TRANSMEM 330 350 POTENTIAL.
 FT SEQUENCE 397 AA; 42421 MW; 678DC6E2A472BF4 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 397;
 Best Local Similarity 75.0%; Pred. No. 25;
 Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPMPWR 10
 I I I I I
 Db 351 RWPMPWR 358

RESULT 12
 YDW6_SCHPO STANDARD; PRT; 535 AA.
 AC Q13912;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, last sequence update)
 DT 15-JUL-1998 (Rel. 36, last annotation update)
 DE HYPOTHEICAL 60.1 KDA PROTEIN C23C11.06C IN CHROMOSOME I.
 GN SPAC23C11.06C.
 OS Schizosaccharomyces pombe (Fission yeast).
 CC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
 CC Schizosaccharomycetales; Schizosaccharomycetaceae;
 CC Schizosaccharomycetes.
 CC NCI_TaxID=4896;
 RX SEQUENCE FROM N.A.
 RP STRAIN-972;
 RC Brown D., Churcher C.M., Barrell B.G., Rajandream M.A., Wood V.;
 RA Submitted (AUG-1997) to the EMBL/Genbank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -----
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 CC -----
 DR EMBL: 298559; CAB1159.1; -
 KW Hypothetical protein; Transmembrane.
 FT TRANSMEM 55 75 POTENTIAL.
 FT TRANSMEM 82 102 POTENTIAL.
 FT TRANSMEM 115 135 POTENTIAL.
 FT TRANSMEM 143 163 POTENTIAL.
 FT TRANSMEM 201 221 POTENTIAL.
 FT TRANSMEM 346 366 POTENTIAL.
 FT TRANSMEM 60124 MW; A6AE149AA2929E2 CRC64;
 SO SEQUENCE

Query Match 51.2%; Score 44; DB 1; Length 535;
 Best Local Similarity 50.0%; Pred. No. 32;
 Matches 6; Conservative 1; Mismatches 1; Indels 4; Gaps 1;

OY 4 WPMW---WPMWR 1
 Db 183 WSMSPSTWPMRQ 194

RESULT 13
 MML4_MYCTU STANDARD; PRT; 967 AA.
 AC 053735;
 DT 30-MAY-2000 (Rel. 39, Created)
 DT 30-MAY-2000 (Rel. 39, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML4.
 GN MML4 OR RV0450C OR MT0466 OR MT037.14C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OC NCBI_TaxID=1773;
 [1]
 RN SEQUENCE FROM N.A.
 RP STRAIN=H37RV;
 RC MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 Gordon S.V., Eigemeier K., Gas S., Barry C.E. III, Tekala F.,
 Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 Davies R., Devlin K., Feltham T., Gentles S., Hamlin N., Holroyd S.,
 Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
 Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
 Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 [2]
 RN SEQUENCE FROM N.A.
 RP STRAIN=CDC 1551 / Oshkosh;
 RC Fleisichmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
 Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
 Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
 Bishai W.;
 RA Delcher A., Uterback T., Weidman J., Khouri H., Gill J., Mikula A.,
 Bishai W.;
 RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RT laboratory strains."
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML FAMILY.
 CC -----
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 CC -----
 DR EMBL: AL021932; CA17407.1; -
 DR EMBL: AE006949; AAK4689.1; -
 DR TIGR: MT0466; -
 KW Tuberculosis; RV0450C;
 FT TRANSMEM 26 46 POTENTIAL.
 FT TRANSMEM 210 230 POTENTIAL.
 FT TRANSMEM 242 262 POTENTIAL.
 FT TRANSMEM 303 323 POTENTIAL.
 FT TRANSMEM 333 353 POTENTIAL.
 FT TRANSMEM 384 404 POTENTIAL.
 FT TRANSMEM 769 789 POTENTIAL.
 FT TRANSMEM 793 813 POTENTIAL.
 FT TRANSMEM 821 841 POTENTIAL.
 FT TRANSMEM 875 895 POTENTIAL.
 FT TRANSMEM 896 916 POTENTIAL.
 FT TRANSMEM 105234 MW; 6301014031480484 CRC64;
 SO SEQUENCE

Query Match 51.2%; Score 44; DB 1; Length 967;
 Best Local Similarity 75.0%; Pred. No. 55;
 Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPMPWR 10
 Db 930 RWPMPWR 937

RESULT 14
 MML2_MYCTU STANDARD; PRT; 968 AA.
 AC 011171;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 20-AUG-2001 (Rel. 40, Last annotation update)
 DE PUTATIVE MEMBRANE PROTEIN MML2.
 GN MML2 OR RV0507 OR MT0528 OR MT0209.34.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 OC NCBI_TaxID=1773;
 [1]
 RN SEQUENCE FROM N.A.
 RP STRAIN=H37RV;
 RC MEDLINE=98295987; PubMed=9634230;
 RA Cole S.T., Brosch R., Parkhill J., Garnier T., Churcher C., Harris D.,
 Gordon S.V., Eigemeier K., Gas S., Barry C.E. III, Tekala F.,
 Badcock K., Basham D., Brown D., Chillingworth T., Connor R.,
 Davies R., Devlin K., Feltham T., Gentles S., Hamlin N., Holroyd S.,
 Hornsby T., Jagels K., Krogh A., McLean J., Moule S., Murphy L.,
 Oliver S., Osborne J., Quail M.A., Rajandream M.A., Rogers J.,
 Rutter S., Seeger K., Skelton S., Squares S., Squares R.,
 Sulston J.E., Taylor K., Whitehead S., Barrell B.G.;
 RA "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence."
 RL Nature 393:537-544(1998).
 [2]
 RN SEQUENCE FROM N.A.
 RP STRAIN=CDC 1551 / Oshkosh;
 RC Fleisichmann R.D., Alland D., Eisen J.A., Carpenter L., White O.,
 Peterson J., Deboy R., Dodson R., Gwinn M.L., Haft D., Hickey E.,
 Kolony J.F., Nelson W.C., Umayam L.A., Ermolaeva M.D., Salzberg S.L.,
 Bishai W.;
 RA Delcher A., Uterback T., Weidman J., Khouri H., Gill J., Mikula A.,
 Bishai W.;
 RA "Whole genome comparison of Mycobacterium tuberculosis clinical and
 RT laboratory strains."
 RL Submitted (APR-2001) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE MML FAMILY.

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DR EMBL: Z77162; CAB00933.1; -
DR EMBL: AE006953; AAK44751.1; -
DR TIGR: MT0528; -
DR TubercuList: RV0507; -
KW Hypothetical protein: Transmembrane: Complete proteome.
FT TRANSMEM 22 42 POTENTIAL.
FT TRANSMEM 204 224 POTENTIAL.
FT TRANSMEM 245 265 POTENTIAL.
FT TRANSMEM 297 317 POTENTIAL.
FT TRANSMEM 328 348 POTENTIAL.
FT TRANSMEM 378 398 POTENTIAL.
FT TRANSMEM 763 783 POTENTIAL.
FT TRANSMEM 787 807 POTENTIAL.
FT TRANSMEM 815 835 POTENTIAL.
FT TRANSMEM 866 886 POTENTIAL.
FT TRANSMEM 891 911 POTENTIAL.
FT CONFLICT 426 426 R -> H (IN REF. 2).
FT CONFLICT 656 656 E -> A (IN REF. 2).
SQ SEQUENCE 968 AA; 106201 MW; B68AB9B78164EDC0 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 968;
Best Local Similarity 75.0%; Pred. No. 55;
Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPWPWR 10
 11111
DB 924 RWPWPWR 931

RESULT 15
SX13_MOUSE STANDARD; PRT; 984 AA.
ID SX13_MOUSE
AC 004891;
DT 01-JUN-1994 (Rel. 29, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 20-AUG-2001 (Rel. 40, Last annotation update)
DE SOX-13 PROTEIN.
GN SOX13 OR SOX-13.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A., AND FUNCTION (ISOFORM 1).
RC TISSUE=Embryo;
RX MEDLINE=98083175; PubMed=9421502;
RA Lamers W., Koorver W., Oving E., Wilson A., Wagenaar G., Markman M.,
RT "High expression of the HMG box factor sox-13 in arterial walls during
RT embryonic development."
RT Nucleic Acids Res. 26:469-476(1998).
RN [2]
RP SEQUENCE FROM N.A., AND FUNCTION (ISOFORM 2).
RC TISSUE=Embryo;
RX MEDLINE=98201614; PubMed=9524265;
RA Kido S., Hiracka Y., Ogawa M., Sakai Y., Yoshimura Y., Aiso S.,
RT "Cloning and characterization of mouse msox13 cDNA."
RT Gene 208:201-206(1998).
RN [3]
RP SEQUENCE OF 405-460 FROM N.A.
RX MEDLINE=93181275; PubMed=8441686;
RA Wright E.M., Snopce B., Koopman P.,
RT "Seven new members of the Sox gene family expressed during mouse

RT development."
RL Nucleic Acids Res. 21:744-744(1993).
CC -1- FUNCTION: BINDS TO THE SEQUENCE 5'-AACAAAT-3'.
CC -1- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS; ISOFORM 1 (SHOWN HERE) AND
CC ISOFORM 2; MAY BE PRODUCED BY ALTERNATIVE SPLICING.
CC -1- TISSUE SPECIFICITY: IN THE EMBRYO, HIGH LEVELS OF EXPRESSION ARE
CC FOUND IN THE ARTERIAL WALLS AT 13.5 DAYS POST COITUM (DPC). LOW
CC LEVELS ARE FOUND IN THE INNER EAR AT 13.5 DPC AND IN SOME CELLS IN
CC THE THYMUS AT 16.5 DPC. EXPRESSED IN THE TRACHEAL EPITHELIUM BELOW
CC THE VOCAL CORD AND IN THE HAIR FOLLICLES AT 18 DPC.
CC -1- SIMILARITY: CONTAINS 1 HMG BOX.

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DR EMBL: AJ000740; CAA04278.1; -
DR EMBL: AB006329; BAA25786.1; -
DR EMBL: Z18962; CAA79487.1; -
DR PIR: S30241; S30241.
DR HSSP: Q05066; 1HRZ.
DR MGP: MGI:98361; Sox13.
DR InterPro: IPR000910; HMG_12_box.
DR Pfam: PF00505; HMG_box; 2.
DR SMART: SM00398; HMG; 1.
DR DNA-binding: Nuclear protein; Alternative splicing.
DR DOMAIN 159 195
FT DNA-BIND 397 465
FT VARSPLIC 495 519
FT VARSPLIC 603 609
FT VARSPLIC 610 984
FT CONFLICT 35 35
FT CONFLICT 41 42
FT CONFLICT 195 195
SQ SEQUENCE 984 AA; 108897 MW; 7F5506EDADEB98C5 CRC64;

Query Match 51.2%; Score 44; DB 1; Length 984;
Best Local Similarity 42.9%; Pred. No. 56;
Matches 6; Conservative 0; Mismatches 0; Indels 8; Gaps 1;

OY 4 WPMW-----PW 9
 11111
DB 686 WPMWTKLAEFSPW 699

Search completed: January 4, 2002, 08:47:48
Job time: 406 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: January 4, 2002, 08:41:32 ; Search time 27.18 seconds
(without alignments)
33.631 Million cell updates/sec

Title: US-09-444-281-36
Perfect score: 86
Sequence: 1 ILRMPWMPWRRK 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 219241 seqs, 76174552 residues

Total number of hits satisfying chosen parameters: 219241

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

1: PIR-68:*
2: PIR1:*
3: PIR2:*
4: PIR3:*
5: PIR4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	70	81.6	144	1 JCI1222	Indolicidin precursor
2	53	61.6	1173	1 VG1HMC	E2 glycoprotein pr
3	51	59.3	299	2 T12505	hypothetical prote
4	50	58.1	111	2 T29295	hypothetical prote
5	49	57.0	198	1 J70751	ferredoxin-NADP+
6	48.5	56.4	114	2 T36208	hypothetical prote
7	47	54.7	248	2 S23449	NADH oxidase (H2O2
8	47	54.7	253	2 G70715	hypothetical prote
9	47	54.7	276	2 B83161	probable short-cha
10	47	54.7	715	2 B70741	probable moef prot
11	47	54.7	1411	2 T48529	hypothetical prote
12	46	53.5	728	2 T51071	related to trifa pr
13	45.5	52.9	505	2 A39128	anthranilate synth
14	45	52.3	196	2 S55483	modulator of drug
15	45	52.3	273	2 F82646	monofunctional dio
16	45	52.3	412	2 A83604	probable MFS trans
17	45	52.3	448	2 H72376	hypothetical prote
18	45	52.3	1108	2 A48508	cyclic-nucleotide
19	44	51.2	257	2 S70177	yife protein - Yer
20	44	51.2	361	2 A36669	galactoside 3(4)-L
21	44	51.2	397	2 B70763	probable membrane
22	44	51.2	535	2 T38244	hypothetical prote
23	44	51.2	621	2 S37664	peplomeric polypyr
24	44	51.2	630	2 S37663	peplomeric polypyr
25	44	51.2	967	2 C70831	probable mmp12 pro
26	44	51.2	968	2 F70746	probable mmp12 pro
27	44	51.2	968	2 T00322	hypothetical prote
28	44	51.2	1154	1 VG1HIB	E2 glycoprotein pr
29	44	51.2	1162	1 VG1HAK	E2 glycoprotein pr

30	44	51.2	1162	2 S07421	E2 glycoprotein pr
31	44	51.2	1162	2 S14939	E2 glycoprotein pr
32	44	51.2	1162	2 S14940	E2 glycoprotein pr
33	43.5	50.6	1529	2 A59189	ATP-binding caset
34	43	50.0	51	2 S23291	light-harvesting p
35	43	50.0	192	2 H86543	hypothetical prote
36	43	50.0	192	2 D72081	conserved hypotet
37	43	50.0	236	2 J00606	arylesterase (EC 3
38	43	50.0	250	2 A83506	probable cobalamin
39	43	50.0	278	2 T46458	hypothetical prote
40	43	50.0	298	2 B72492	hypothetical prote
41	43	50.0	646	2 H82555	c-type cytochrome
42	43	50.0	711	2 C40046	antibiotic transpo
43	43	50.0	738	2 F96701	hypothetical prote
44	43	50.0	958	2 A70634	probable mmp12 pro
45	43	50.0	1112	2 S70522	cyclic nucleotide

ALIGNMENTS

RESULT 1
JCI1222
Indolicidin precursor - bovine
N:Alternate names: antimicrobial peptide
C:Species: Bos primigenius taurus (cattle)
C>Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
C:Accession: JCI1222; A42387; S25664
Ridel Sal, G.; Storici, P.; Schneider, C.; Romeo, D.; Zanetti, M.
Biochem. Biophys. Res. Commun. 187, 467-472, 1992
A:Title: cDNA cloning of the neutrophil bactericidal peptide indolicidin.
A:Reference number: JCI1222; MUID:92392368
A:Accession: JCI1222
A:Molecule type: mRNA
A:Residues: 1-144 <SAL>
A:Cross-references: EMBL:X67340; NID:9462; PID:CAA47755.1; PID:9463
A:Experimental source: Bone marrow
R:Seidel, M.E.; Novotny, M.J.; Morris, W.L.; Tang, Y.Q.; Smith, W.; Cullor, J.S.
J. Biol. Chem. 267, 4292-4295, 1992
A:Title: Indolicidin, a novel bactericidal tridecapeptide amide from neutrophils.
A:Reference number: A42387; MUID:92165771
A:Accession: A42387
A:Residues: 131-143 <SEL>
A:Molecule type: protein
A:Residues: 131-143 <SEL>
A:Experimental source: neutrophils
A>Note: sequence extracted from NCBI backbone (NCBI:83840)
C:Superfamily: cathelin; cystatin homology
C:Keywords: amidated carboxyl end
F:1-29/Domain: signal sequence #status predicted <SIG>
F:32-129/Domain: cystatin homology <CVS>
F:30-130/Domain: propeptide #status predicted <PRO>
F:131-143/Product: indolicidin #status experimental <MAT>
F:143/Modified site: amidated carboxyl end (Arg) (amide in mature form from following

Query Match 81.4% Score 70; DB 1; Length 144;
Best Local Similarity 88.9% Pred. No. 0.011;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 3 RMPWMPWRR 11
DB 135 RMPWMPWRR 143

RESULT 2
VG1HMC
E2 glycoprotein precursor - human coronavirus (strain 229E)
N:Alternate names: peplomer glycoprotein; S glycoprotein; spike glycoprotein
C:Species: human coronavirus
A>Note: host Homo sapiens (man)
C>Date: 31-Dec-1991 #sequence_revision 31-Dec-1991 #text_change 16-Jun-2000
C:Accession: A34766; S05460
R:Raabe, T.; Schelle-Prinz, B.; Siddell, S.G.

J. Gen. Virol. 71, 1065-1073, 1990
 A:Title: Nucleotide sequence of the gene encoding the spike glycoprotein of human corona
 A:Reference number: A34766; MUID:90264837
 A:Accession: A34766
 A:Molecule type: mRNA
 A:Residues: 1-1173 <RA>
 A:Cross-references: EMBL:X16816; NID:958926; PTDN:CAA34723.1; PID:958927
 A:Experimental source: strain 229E
 R:Raabe, T.; Sidde11, S.
 Nucleic Acids Res. 17, 6387, 1989
 A:Title: Nucleotide sequence of the human coronavirus HCV 229E mRNA 4 and mRNA 5 unique
 A:Reference number: A34038; MUID:89366667
 A:Accession: S05460
 A:Status: translation not shown
 A:Molecule type: mRNA
 A:Residues: 1159-1173 <RA>
 A:Cross-references: EMBL:X15654; NID:958921; PTDN:CAA33680.1; PID:91334827
 C:Superfamily: coronavirus E2 glycoprotein
 C:Keywords: glycoprotein; transmembrane protein
 F:1-15/Domain: signal sequence #status predicted <SIG>
 F:16-1173/Product: E2 glycoprotein #status predicted <MAT>
 F:1116-1138/Domain: transmembrane #status predicted <TM>
 F:23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,
 5;23,62,98,147,171,176,220,243,326,333,440,464,518,538,542,568,581,587,663,671,930,1015,

Query Match 61.6%; Score 53; DB 1; Length 1173;
 Best Local Similarity 62.5%; Pred. No. 12;
 Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 LRMPWPMW 9
 Db 1112 IKMPWPMW 1119

RESULT 3
 T12505
 hypothetical protein DKFZp434C192.1 - human (fragment)
 C:Species: Homo sapiens (man)
 C:Date: 23-Jul-1999 #sequence_revision 23-Jul-1999 #text_change 23-Jul-1999
 C:Accession: T12505
 R:Ansorge, W.; Wilkner, U.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
 submitted to the Protein Sequence Database, June 1999
 A:Reference number: 217527
 A:Accession: T12505
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-299 <ANS>
 A:Cross-references: EMBL:A1096753
 A:Experimental source: adult testis; clone DKFZp434C192
 C:Genetics:
 A:Note: DKFZp434C192.1

Query Match 59.3%; Score 51; DB 2; Length 299;
 Best Local Similarity 85.7%; Pred. No. 6.1;
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 5 PMPMPMR 11
 Db 37 PMPMPMR 43

RESULT 4
 T29295
 hypothetical protein C50F7.8 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 15-Oct-1999
 C:Accession: T29295
 R:Johnson, D.; Steilly, L.
 submitted to the EMBL Data Library, November 1995
 A:Description: The sequence of C. elegans cosmid C50F7.
 A:Reference number: Z20601
 A:Accession: T29295

A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-111 <IOH>
 A:Cross-references: EMBL:U41557; PTDN:AAA83303.1; CESP:C50F7.8
 C:Genetics:
 A:Gene: CESP:C50F7.8

Query Match 58.1%; Score 50; DB 2; Length 111;
 Best Local Similarity 54.5%; Pred. No. 3.2;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

OY 1 LRMPWPMW 11
 Db 12 VMWPMWPGCR 22

RESULT 5
 JF0751
 ferredoxin-NADP+ reductase (EC 1.18.1.2), long form precursor - bovine
 N:Alternate names: adrenodoxin reductase
 C:Species: Bos primigenius taurus (cattle)
 C:Date: 14-Jul-1994 #sequence_revision 18-Oct-1996 #text_change 16-Jun-2000
 C:Accession: JF0751; JF0079; JS0390; S03558; PS0003; A29604; S52100
 R:Takata, Y.; Sagara, Y.; Kono, A.; Sekimizu, K.; Horiiuchi, T.
 Biol. Pharm. Bull. 16, 1200-1206, 1993
 A:Title: Gene structure of bovine adrenodoxin reductase.
 A:Reference number: JF0751; MUID:94177140
 A:Accession: JF0751
 A:Molecule type: DNA
 A:Residues: 1-498 <RA>
 A:Cross-references: GB:D83475; NID:q1199916; PTDN:BAH11921.1; PID:94521308
 A:Experimental source: adrenal cortex
 A:Note: the authors translated the codon GTC for residue 205 as Gly
 R:Sagara, Y.; Takata, Y.; Miyata, T.; Hara, T.; Horiiuchi, T.
 J. Biochem. 102, 1333-1336, 1987
 A:Title: Cloning and sequence analysis of adrenodoxin reductase cDNA from bovine adre
 A:Reference number: JF0079; MUID:88198050
 A:Accession: JF0079
 A:Molecule type: mRNA
 A:Residues: 1-204,211-498 <SAG>
 A:Cross-references: GB:D00211; NID:q217433; PTDN:BAH00150.1; PID:q217434
 A:Note: the deduced sequence is partially confirmed by amino acid sequencing of 15 is
 R:Sagara, Y.
 submitted to DDBJ, September 1989
 A:Reference number: JS0390
 A:Contents: revision, insertion of residues 205-210
 A:Accession: JS0390
 A:Molecule type: mRNA
 A:Residues: 56-498 <SAG>
 R:Hankoglu, I.; Gutfinger, T.
 Eur. J. Biochem. 180, 479-484, 1989
 A:Title: cDNA sequence of adrenodoxin reductase. Identification of NADP-binding sites
 A:Reference number: S03558; MUID:89170752
 A:Accession: S03558
 A:Molecule type: mRNA
 A:Residues: 135-204,211-498 <HAN>
 A:Cross-references: EMBL:X13736; NID:9665; PTDN:CAA32002.1; PID:9833776
 A:Note: 405-Ser was also found
 R:Hamamoto, I.; Kurokouchi, K.; Tanaka, S.; Ichikawa, Y.
 Biochim. Biophys. Acta 953, 207-213, 1988
 A:Title: Adrenoferritin-binding peptide of NADPH-adrenoferritin reductase.
 A:Reference number: PS0003; MUID:88184054
 A:Accession: PS0003
 A:Molecule type: protein
 A:Residues: 33-41, 'S', '43-62:260-283, 'TM', '496-498 <HAM>
 A:Note: a cyanogen bromide peptide binds to adrenoferritin
 R:Nonaka, Y.; Murakami, H.; Yanuski, Y.; Kuramitsu, S.; Kagamiyama, H.; Yamano, T.;
 Biochem. Biophys. Res. Commun. 145, 1239-1247, 1987
 A:Title: Molecular cloning and sequence analysis of full-length cDNA for adre
 A:Reference number: A29604; MUID:87270696
 A:Accession: A29604
 A:Molecule type: mRNA

A:Residues: 1-76, 'R', '78-80, 'VMALTPRSRL', '95-123, 'RVYRLT', '129-204, '211-273, 'R', '275-322, 'A:Cross-references: GB:M17029; NID:g162628; PIDN:AAA30362.1; PID:g162629
A:Experimental source: adrenal cortex
R:Warburton, R.J.; Seybert, D.W.
Biochim. Biophys. Acta 1246, 39-46, 1995
A:Title: Structural and functional characterization of bovine adrenodoxin reductase by 1
A:Reference number: S52100; MUID:95110846
A:Accession: S52100
A:Status: preliminary
A:Molecule type: protein
A:Residues: 'X', '34-41, 'X', '43-48, 'X', '50-51, '304-306, 'X', '308-309, 'X', '311-326 <MAR>
C:Comment: Ferredoxin-NADP+ reductase is localized in the matrix of adrenal cortex mito
ferredoxin-NADP+ reductase, adrenodoxin and two forms of cytochrome P-450.
C:Genetics:
A:Introns: 27/1; 59/3; 91/3; 132/3; 170/3; 204/3; 246/3; 275/1; 341/3; 399/1; 456/1
C:Function:
A:Description: catalyzes the reversible reduction of NADP+ by reduced ferredoxin or red
A:Superfamily: human ferredoxin-NADP+ reductase
C:Keywords: alternative splicing; flavoprotein; mitochondrion; monomer; NADP; oxidoreduc
F:1-32/Domain: transit peptide (mitochondrion) #status predicted <SIG>
F:33-498/Product: ferredoxin-NADP+ reductase, long form #status predicted <MAT>
F:33-204, 211-498/Product: ferredoxin-NADP+ reductase, short form #status experimental <
F:40-70/Region: beta-alpha-beta FAD nucleotide-binding fold
F:180-190/Region: NADP binding #status predicted
F:281/Binding site: substrate (Lys) #status experimental

Query Match 57.0%; Score 49; DB 1; Length 498;
Best Local Similarity 83.3%; Pred. No. 18;
Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 4 PWMPPW 9
1 1111
DB 6 WRMPW 11

RESULT 6
136208
hypothetical protein SCE36.09 - Streptomyces coelicolor
C:Species: Streptomyces coelicolor
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Dec-1999
C:Accession: J36208
R:Oliver, K.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
submitted to the EMBL Data Library, May 1999
A:Reference number: Z21601
A:Accession: J36208
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-114 <OLI>
A:Cross-references: EMBL:AL049763; PIDN:CAB42078.1; GSPDB:GN00070; SCOEDB:SCE36.09
A:Experimental source: strain A3(2)
C:Genetics:
A:Gene: SCOEDB:SCE36.09

Query Match 56.4%; Score 48.5; DB 2; Length 114;
Best Local Similarity 80.0%; Pred. No. 5.1;
Matches 8; Conservative 0; Mismatches 1; Indels 1; Gaps 1;

QY 3 RW-PWMPWR 11
1 111111
DB 103 RWRPMPWR 112

RESULT 7
S23449
NADH oxidase (H2O2-forming) (EC 1.6.1.1) - Thermus aquaticus
C:Species: Thermus aquaticus
C:Date: 22-Jan-1993 #sequence_revision 22-Jan-1993 #text_change 23-Mar-1993
C:Accession: S23449; S24556
R:Park, H.U.; Kreutzer, R.; Reiser, C.O.A.; Sprinzl, M.
Eur. J. Biochem. 205, 875-879, 1992
A:Title: Molecular cloning and nucleotide sequence of the gene encoding a H(2)O(2)-formi

A:Reference number: S23449; MUID:92249331
A:Accession: S23449
A:Molecule type: DNA
A:Residues: 1-248 <PAR>
A:Cross-references: EMBL:X60110
A:Accession: S24556
A:Molecule type: protein
A:Residues: 1-32 <PAR1>
C:Genetics:
A:Gene: nox
C:Keywords: NAD; oxidoreductase
F:1-248/Product: NADH oxidase (H2O2-forming) #status experimental <MAT>

Query Match 54.7%; Score 47; DB 2; Length 248;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 5 PWMPPW 9
1 1111
DB 179 PWMPW 183

RESULT 8
G70715
hypothetical protein RV0945 - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 20-Jun-2000
C:Accession: G70715
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
; Connor, R.; Davies, R.; Devlin, K.; Fellwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
Nature 393, 537-544, 1998
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
A:Reference number: A70500; MUID:98295987
A:Accession: G70715
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-253 <COL>
A:Cross-references: GB:279700; GB:AL133456; NID:g3261628; PIDN:CAB02005.1; PID:g15242
A:Experimental source: strain H37RV
C:Genetics:
A:Gene: RV0945
C:Superfamily: ribitol dehydrogenase; short-chain alcohol dehydrogenase homology
F:8-190/Domain: short-chain alcohol dehydrogenase homology <SADH>

Query Match 54.7%; Score 47; DB 2; Length 253;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 PWMPPW 9
1 1111
DB 230 PWMPW 234

RESULT 9
B83161
probable short-chain dehydrogenase PA3883 [Imported] - Pseudomonas aeruginosa (strain
C:Species: Pseudomonas aeruginosa
C:Date: 15-Sep-2000 #sequence_revision 15-Sep-2000 #text_change 31-Dec-2000
C:Accession: B83161
R:Stover, C.K.; Pham, X.O.; Erwin, A.L.; Mizoguchi, S.D.; Warrenner, P.; Hickey, M.J.;
adman, S.; Yuan, Y.; Brody, L.L.; Coulter, S.N.; Folger, K.R.; Kas, A.; Lardig, K.; L
; Lory, S.; Olson, M.V.
Nature 406, 959-964, 2000
A:Title: Complete genome sequence of Pseudomonas aeruginosa PA01, an opportunistic pa
A:Reference number: A82950; MUID:20437337
A:Accession: B83161
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-276 <STO>

A:Cross-references: GB:AE004805; GB:AE004091; NID:g950055; PIDN:AG07270.1; GSPDB:GN001
 A:Experimental source: straln PA01
 C:Genetics:
 A:Gene: PA3883
 C:Superfamily: ribitol dehydrogenase; short-chain alcohol dehydrogenase homology

Query Match 54.7%; Score 47; DB 2; Length 276;
 Best Local Similarity 60.0%; Pred. No. 18;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 3 RWPMPWRRK 12
 Db 197 RWPMPWRRK 206

RESULT 10
 B70741
 probable moey protein - Mycobacterium tuberculosis (strain H37RV)

C:Species: Mycobacterium tuberculosis
 C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 22-Oct-1999
 C:Accession: B70741
 R:Coile, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Felwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 Nature 393, 537-544, 1998
 A:Authors: Squires, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome
 A:Reference number: A70500; MUID:98295987
 A:Accession: B70741
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-715 <COL>
 A:Cross-references: GB:Z75555; GB:AL123456; NID:g3261608; PIDN:CA99988.1; PID:e250356;
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: moey

Query Match 54.7%; Score 47; DB 2; Length 715;
 Best Local Similarity 66.7%; Pred. No. 46;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 3 RWPMPWRRK 11
 Db 65 RWPMPWRRK 73

RESULT 11
 T48529
 hypothetical protein T22P22.90 - Arabidopsis thaliana
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 C:Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 20-Apr-2000
 C:Accession: T48529
 R:Bevan, M.; Hilbert, H.; Braun, M.; Holzer, E.; Brandt, A.; Duesterhoft, A.; Bancroft, submitted to the Protein Sequence Database, April 2000
 A:Reference number: Z24490
 A:Accession: T48529
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-1411 <BEV>
 A:Cross-references: EMBL:AL163814
 A:Experimental source: cultivar Columbia; BAC clone T22P22
 C:Genetics:
 A:Map position: 5
 A:Introns: 281/2; 320/1; 389/3; 429/3; 473/3; 515/3; 534/2; 567/3; 602/1; 669/1; 776/2;
 A:Note: T22P22.90

Query Match 54.7%; Score 47; DB 2; Length 1411;
 Best Local Similarity 63.6%; Pred. No. 87;
 Matches 7; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 2 LRMPMPWRRK 12
 Db 1013 LRMPMPWRRK 1023

RESULT 12
 T51071
 related to trfA protein [imported] - Neurospora crassa
 N:Alternate names: protein B2A19.50
 C:Species: Neurospora crassa
 C:Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
 C:Accession: T51071
 R:Schulte, U.; Algen, V.; Hohnsels, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatu submitted to the Protein Sequence Database, July 2000
 A:Reference number: Z25286
 A:Accession: T51071
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-728 <SCH>
 A:Cross-references: EMBL:AL390092; GSPDB:GN00116; NCSP:B2A19.50
 A:Experimental source: BAC clone B2A19; strain OR74A
 C:Genetics:
 A:Gene: NCSP:B2A19.50
 A:Map position: 6
 A:Introns: 26/1; 119/2

Query Match 53.5%; Score 46; DB 2; Length 728;
 Best Local Similarity 58.3%; Pred. No. 62;
 Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

OY 1 LRMPMPWRRK 12
 Db 11 LRMPMPWRRK 22

RESULT 13
 A39128
 anthranilate synthase (EC 4.1.3.27) component I [validated] - Pseudomonas syringae pv
 N:Alternate names: anthranilate synthase alpha chain
 C:Species: Pseudomonas syringae pv. savastanoi
 C:Date: 27-Nov-1991 #sequence_revision 27-Nov-1991 #text_change 17-Mar-2000
 C:Accession: A39128
 R:da Costa, E.; Silva, O.; Kosuge, T.
 J. Bacteriol. 173, 463-471, 1991
 A:Title: Molecular characterization and expression analysis of the anthranilate synth
 A:Reference number: A39128; MUID:91100331
 A:Accession: A39128
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-505 <DAC>
 A:Cross-references: GB:M55911
 C:Genetics:
 A:Gene: trpE
 C:Complex: heterotrimer; two component I chains, two component II chains
 C:Function: <ANT>
 A:Description: EC 4.1.3.27 [validated; MUID:90130325]
 A:Pathway: tryptophan biosynthesis
 A:Note: first step
 C:Function: <COMI>
 A:Description: EC 4.1.3.27 [validated; MUID:91100331]
 A:Note: expression of trpE seems to be independent of the concentration of tryptophan
 C:Superfamily: anthranilate synthase component I
 C:Keywords: carbon-carbon lyase; oxo-acid-lyase; tryptophan biosynthesis

Query Match 52.9%; Score 45.5; DB 2; Length 505;
 Best Local Similarity 28.0%; Pred. No. 51;
 Matches 7; Conservative 2; Mismatches 3; Indels 13; Gaps 1;

OY 1 LRMPMPWRRK 12
 Db 467 LRMPMPWRRK 491

RESULT 14

S55483

modulator of drug activity homolog, fission yeast (Schizosaccharomyces pombe)

C:Species: Schizosaccharomyces pombe

C:Date: 01-Aug-1995 #sequence_revision 01-Sep-1995 #text_change 10-Dec-1999

C:Accession: S55483; T38969

R:Connor, R.; Churcher, C.M. submitted to the EMBL Data Library, May 1995

A:Reference number: S55479

A:Accession: S55483

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-196 <CON>

A:Cross-references: EMBL:Z49811; NID:9854599; PIDN:CAA89955.1; PID:9854604

R:Connor, R.; Churcher, C.M.; Barrall, B.G.; Rajandream, M.A.; Walsh, S.V. submitted to the EMBL Data Library, May 1995

A:Reference number: Z21821

A:Accession: T38969

A:Status: preliminary; translated from GR/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-196 <CO2>

A:Cross-references: EMBL:Z49811; PIDN:CAA89955.1; GSPDB:GN00066; SPDB:SPAC5H10.05c

A:Experimental source: strain 972h-; cosmid c5H10

C:Genetics:

A:Gene: SPDB:SPAC5H10.05c

A:Map position: 1

C:Superfamily: MAD(P)H dehydrogenase (quinone) 2

Query Match 52.3%; Score 45; DB 2; Length 196;

Best Local Similarity 50.0%; Pred. No. 24;

Matches 8; Conservative 2; Mismatches 2; Indels 4; Gaps 2;

OY 1 ILRWP-WM---PWRK 12

1 :||| 11 : 1

Db 63 IYQMPGMMGCTPWKLK 78

RESULT 15

F82646

monofunctional biosynthetic,peptidoglycan transglycosylase XF1715 [Imported] - Xylella

C:Species: Xylella fastidiosa

C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 20-Aug-2000

C:Accession: F82646

R:anonymous; The Xylella fastidiosa Consortium of the Organization for Nucleotide Sequen

Nature 406, 151-157, 2000

A:Title: The genome sequence of the plant pathogen Xylella fastidiosa.

A:Reference number: A82515; MUID:20365717

A>Note: for a complete list of authors see reference number A59328 below

A:Accession: F82646

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-273 <SIM>

A:Cross-references: GB:AE003995; GB:AE003849; NID:9106775; PIDN:AAF84524.1; GSPDB:GN001

A:Experimental source: strain 9a5c

R:Simpson, A.J.G.; Reinach, F.C.; Arruda, P.; Abreu, F.A.; Agencio, M.; Alvarenga, R.; A

Briones, M.R.S.; Bueno, M.R.P.; Camargo, A.A.; Camargo, L.E.A.; Carraro, D.M.; Carreir, H

as-Neto, E.; Docena, C.; El-Dorri, H.; Facinanci, A.P.; Ferreira, A.J.S.

A:Authors: Ferreira, V.C.A.; Ferro, J.A.; Fraga, J.S.; Franca, S.C.; Franco, M.C.; Frohm

J.D.; Junqueira, M.L.; Kemper, E.L.; Kitajima, J.P.; Krieger, J.E.; Kuramae, E.E.; Laig

chado, M.A.; Madeira, A.M.B.N.; Madeira, H.M.F.; Marino, C.L.; Marques, M.V.; Martins, E

A:Authors: Martins, E.M.F.; Matsukuma, A.Y.; Menck, C.F.M.; Miracca, E.C.; Miyaki, C.Y.;

F.G.; Nunes, L.R.; Oliveira, M.A.; de Oliveira, M.C.; de Oliveira, R.C.; Palmieri, D.A

Rodrigues, V.; Rosa, A.J. de M.; de Rosa Jr., V.E.; de Sa, R.G.; Santelli, R.V.; Sawasak

A:Authors: da Silva, A.C.R.; da Silva, F.R.; da Silva, A.M.; Silva Jr., W.A.; da Silveir

M.; Tsubako, M.H.; Vallada, H.; Van Sluys, M.A.; Verjovski-Almeida, S.; Vettore, A.L.; Z

A:Reference number: A59328

A:Contents: annotation

C:Genetics:

A:Gene: XF1715

Query Match 52.3%; Score 45; DB 2; Length 273;

Best Local Similarity 100.0%; Pred. No. 33;

Matches 5; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 RMPW 7

11111

Db 44 RMPW 48

Search completed: January 4, 2002, 08:41:33
Job time: 171 sec

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APPLICANT: Kim, Jeong Hyun
APPLICANT: Hong, Seung-Suh
APPLICANT: Lee, Hyun-Soo
APPLICANT: Samyang Genex Corporation
TITLE OF INVENTION: METHOD FOR MASS PRODUCTION OF
FILE REFERENCE: 6181/0F135
CURRENT APPLICATION NUMBER: US/09/230,180
CURRENT FILING DATE: 1999-03-10
PRIOR APPLICATION NUMBER: PCT/KR98/00132
PRIOR FILING DATE: 1998-05-28
PRIOR APPLICATION NUMBER: KR 13372/1998
PRIOR FILING DATE: 1998-04-09
PRIOR APPLICATION NUMBER: KR 21312/1997
PRIOR FILING DATE: 1997-05-28
NUMBER OF SEQ ID NOS: 36
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 29
LENGTH: 39
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: DNA sequence deduced from Indolicidin peptide
US-09-230-180-29

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x US-09-230-180-29 ..

Align seg 1/1 to: US-09-230-180-29 from: 1 to: 39

3 ArgTrpProtTrpProtTrpArgArg 11
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13 AATGGCGGTGGTGGCGGTGGT 39

seq_name: /cgn2_6/ptodata/2/lna/6A_COMB.seq:US-09-259-741-5

seq_documentation_block:

Sequence 5, Application US/09259741
Patent No. 6033895

GENERAL INFORMATION:

APPLICANT: GARGER, STEPHEN

APPLICANT: HOLTZ, R. BARRY

APPLICANT: MCCULLOCH, MICHAEL

APPLICANT: TURPEN, THOMAS

TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES FROM PLANT

TITLE OF INVENTION: SOURCES

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Howrey & Simon

STREET: 1299 Pennsylvania Avenue N.W.

CITY: Washington

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/259,741

FILING DATE: February 25, 1999

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/037,751
FILING DATE: March 10, 1998
ATTORNEY/AGENT INFORMATION:
NAME: Halluin, Albert P
REGISTRATION NUMBER: 25,277
REFERENCE/DOCKET NUMBER: 00801,0140,US01
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-463-8100
TELEFAX: 650-463-8400
TELEX:
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 6446 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: unknown
MOLECULE TYPE: Genomic RNA
US-09-259-741-5

alignment_scores:
Quality: 70.00 Length: 9
Ratio: 7.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-09-444-281-36 x US-09-259-741-5 ..

Align seg 1/1 to: US-09-259-741-5 from: 1 to: 6446

3 ArgTrpProtTrpProtTrpArgArg 11
:|||||
6213 AAGUGCCUGUGUGCCAGCGCGCA 6239

seq_name: /cgn2_6/ptodata/2/lna/6A_COMB.seq:US-09-037-751-5

seq_documentation_block:

Sequence 5, Application US/09037751
Patent No. 6037456

GENERAL INFORMATION:

APPLICANT: GARGER, STEPHEN

APPLICANT: HOLTZ, R. BARRY

APPLICANT: MCCULLOCH, MICHAEL

APPLICANT: TURPEN, THOMAS

TITLE OF INVENTION: A PROCESS FOR ISOLATING AND
PURIFYING VIRUSES SOLUBLE PROTEINS AND PEPTIDES

TITLE OF INVENTION: FROM PLANT SOURCES

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Howrey & Simon

STREET: 1299 Pennsylvania Avenue N.W.

CITY: Washington

STATE: DC

COUNTRY: USA

ZIP: 20004

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/037,751

FILING DATE: 10-MAR-1998

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER:

FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Halluin, Albert P

REGISTRATION NUMBER: 25,277

REFERENCE/DOCKET NUMBER: 00801,0140,999

TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-463-8109

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: MOLECULE TYPE: Genomic RNA
: SEQUENCE DESCRIPTION: SEQ ID NO: 5
US-09-466-422-5

alignment_scores:
    Quality: 70.00      Length: 9
    Ratio: 7.778      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 88.889

alignment_block:
US-09-444-281-36 x US-09-466-422-5 ..

Align seg 1/1 to: US-09-466-422-5 from: 1 to: 6446

3 ArgTTPrPTPrPrPTPrPTArGArG 11
:::|||||
6213 AAGUGCCUUGGUGGCCAUGGCCCGA 6239

seq_name: /cgn2_6/plodata/2/1na/6A_COMB.seq:US-08-793-035-6

seq_documentation_block:
: Sequence 6, Application US/08793035
: Patent No. 6011201
: GENERAL INFORMATION:
: APPLICANT: Slabas, Antoni R.
: APPLICANT: White, Andrew
: APPLICANT: Chase, Dianne
: APPLICANT: Elborough, Keiran
: APPLICANT: Fentem, Phillip A.
: TITLE OF INVENTION: B-retacyl ACP Reductase Genes From
: TITLE OF INVENTION: Brassica Napus
: NUMBER OF SEQUENCES: 10
: CORRESPONDENCE ADDRESSES:
: ADDRESSEE: Arnold White & Durkee
: STREET: P.O. Box 4433
: CITY: Houston
: STATE: TX
: COUNTRY: US
: ZIP: 77210-4433
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patent Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/793,035
: FILING DATE: 28-JUL-1997
: CLASSIFICATION: B00
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: GB 9414622.2
: FILING DATE: 20-JUL-1994
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: GB95/01678
: FILING DATE: 17-JUL-1995
: ATTORNEY/AGENT INFORMATION:
: NAME: Kammerer, Patricia A.
: REGISTRATION NUMBER: 29,775
: REFERENCE/DOCKET NUMBER: MOST:132
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 713.787.1400
: TELEFAX: 713.787.1440
: INFORMATION FOR SEQ ID NO: 6:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 758 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-08-793-035-6

alignment_scores:
Quality: 61.00      Length: 9

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Ratio: 7.625 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:

US-09-444-281-36 x US-08-793-035-6/rev ..

Align seg 1/1 to reverse of: US-08-793-035-6 from: 1 to: 758

1 l1leuargtrprrprrprrprrp 9
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451 CTCCTGCATGAGTGTGAGCTGG 425

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-020-956-82

seq_documentation_block:

; Sequence 82, Application US/09020956;
; Patent No. 6261562
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, David C.
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FO
; NUMBER OF SEQUENCES: 178
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US709/020,956
; FILING DATE: 09-FEB-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Maki, David J.
; REGISTRATION NUMBER: 210121.427C2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 82:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 383 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-09-020-956-82

alignment_scores:

Quality: 59.00 Length: 8
Ratio: 8.429 Gaps: 0
Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:

US-09-444-281-36 x US-09-020-956-82 ..

Align seg 1/1 to: US-09-020-956-82 from: 1 to: 383

2 leuargtrprrprrprrprrp 9
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155 CTCGCTGCCTGTGAGCTGG 178

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-030-607-82

seq_documentation_block:

; Sequence 82, Application US/09030607
; Patent No. 6262245
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, David C.
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS
; NUMBER OF SEQUENCES: 224
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98104
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/030,607
; FILING DATE: 25-FEB-1998
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Maki, David J.
; REGISTRATION NUMBER: 31,392
; REFERENCE/DOCKET NUMBER: 210121.427C3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 82:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 383 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-09-030-607-82

alignment_scores:

Quality: 59.00 Length: 8
Ratio: 8.429 Gaps: 0
Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:

US-09-444-281-36 x US-09-030-607-82 ..

Align seg 1/1 to: US-09-030-607-82 from: 1 to: 383

2 leuargtrprrprrprrprrp 9
|||||
155 CTCGCTGCCTGTGAGCTGG 178

seq_name: /cgn2_6/ptodata/2/lna/6B_COMB.seq:US-09-030-607-183

seq_documentation_block:

; Sequence 183, Application US/09030607
; Patent No. 6262245
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Dillon, David C.
; TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS
; NUMBER OF SEQUENCES: 224
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED and BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: WA
; COUNTRY: USA


```

: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/030.607
: FILING DATE: 25-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C3
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
: INFORMATION FOR SEQ ID NO: 183:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 384 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
:
: US-09-030-607-183

alignment_scores:
Quality: 59.00 Length: 8
Ratio: 8.429 Gaps: 0
Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:
US-09-444-281-36 x US-09-030-607-183 ..
Align seg 1/1 to: US-09-030-607-183 from: 1 to: 384
2 LeuAgtPrpTrpTrpPrpTrp 9
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156 CTTCGCTGCGCTTGTGTGAGACTGG 179

seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-020-956-73
seq_documentation_block:
: Sequence 73, Application US/09020956
: Patent No. 6261562
: GENERAL INFORMATION:
: APPLICANT: Xu, Jiangchun
: APPLICANT: Dillon, Davin C.
: TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS FO
: NUMBER OF SEQUENCES: 178
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SEED and BERRY LLP
: STREET: 6300 Columbia Center, 701 Fifth Avenue
: CITY: Seattle
: STATE: WA
: COUNTRY: USA
: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/020.956
: FILING DATE: 09-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C2
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
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: INFORMATION FOR SEQ ID NO: 73:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 499 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: ORIGINAL SOURCE:
: ORGANISM: Homo sapiens
:
: US-09-020-956-73

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Percent Similarity: 87.500 Percent Identity: 87.500

alignment_block:
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seq_documentation_block:
: Sequence 73, Application US/09030607
: Patent No. 6262245
: GENERAL INFORMATION:
: APPLICANT: Xu, Jiangchun
: APPLICANT: Dillon, Davin C.
: TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY OF PROSTATE CANCER AND METHODS
: NUMBER OF SEQUENCES: 224
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SEED and BERRY LLP
: STREET: 6300 Columbia Center, 701 Fifth Avenue
: CITY: Seattle
: STATE: WA
: COUNTRY: USA
: ZIP: 98104
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.30
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/030.607
: FILING DATE: 25-FEB-1998
: CLASSIFICATION:
: ATTORNEY/AGENT INFORMATION:
: NAME: Makl, David J.
: REGISTRATION NUMBER: 31,392
: REFERENCE/DOCKET NUMBER: 210121.427C3
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
: INFORMATION FOR SEQ ID NO: 73:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 499 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: ORIGINAL SOURCE:
: ORGANISM: Homo sapiens
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: US-09-030-607-73

alignment_scores:
Quality: 59.00 Length: 8
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Ratio: 8.429 Gaps: 0
Percent Similarity: 87.500 Percent Identity: 87.500

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seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-088-651-3

seq_documentation_block:

Sequence 3, Application US/09088651
Patent No. 6165771

GENERAL INFORMATION:

APPLICANT: BURGESS, NICOLA A.

APPLICANT: CLINKENBEARD, HELEN E.

APPLICANT: SOUTHAN, CHRISTOPHER D.

TITLE OF INVENTION: NOVEL COMPOUNDS

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:

ADDRESSEE: RATNER & PRESTIA

STREET: P.O. BOX 980

CITY: VALLEY Forge

STATE: PA

COUNTRY: USA

ZIP: 19482

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/088,651

FILING DATE: JUNE 1, 1998

CLASSIFICATION:

Prior Application DATA:

APPLICATION NUMBER: GB9712088.5

FILING DATE: 10-JUNE-1997

APPLICATION NUMBER: EP 97308295.1

FILING DATE: 17-OCT-1997

APPLICATION NUMBER: GB 9803650.2

FILING DATE: 20-FEB-1998

ATTORNEY/AGENT INFORMATION:

NAME: PRESTIA, PAUL F.

REGISTRATION NUMBER: 23,031

REFERENCE/DOCKET NUMBER: GH30358

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-407-0700

TELEFAX: 610-407-0701

TELEX: 846169

INFORMATION FOR SEQ. ID NO: 3:

SEQUENCE CHARACTERISTICS:

LENGTH: 683 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-09-088-651-3

alignment_scores:

Quality: 59.00 Length: 11

Ratio: 6.556 Gaps: 0

Percent Similarity: 81.818 Percent Identity: 63.636

alignment_block:

US-09-444-281-36 x US-09-088-651-3 ..

Align seg 1/1 to: US-09-088-651-3 from: 1 to: 683

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||| |||||
139 CTGACGTGGCCCTGTGTGAGCTGAGCAG 171

seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-088-651-1

seq_documentation_block:

Sequence 1, Application US/09088651

Patent No. 6165771

GENERAL INFORMATION:

APPLICANT: BURGESS, NICOLA A.

APPLICANT: CLINKENBEARD, HELEN E.

APPLICANT: SOUTHAN, CHRISTOPHER D.

TITLE OF INVENTION: NOVEL COMPOUNDS

NUMBER OF SEQUENCES: 6

CORRESPONDENCE ADDRESS:

ADDRESSEE: RATNER & PRESTIA

STREET: P.O. BOX 980

CITY: VALLEY Forge

STATE: PA

COUNTRY: USA

ZIP: 19482

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FASTSEQ for Windows Version 2.0

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/088,651

FILING DATE: JUNE 1, 1998

CLASSIFICATION:

Prior Application DATA:

APPLICATION NUMBER: GB9712088.5

FILING DATE: 10-JUNE-1997

APPLICATION NUMBER: EP 97308295.1

FILING DATE: 17-OCT-1997

APPLICATION NUMBER: GB 9803650.2

FILING DATE: 20-FEB-1998

ATTORNEY/AGENT INFORMATION:

NAME: PRESTIA, PAUL F.

REGISTRATION NUMBER: 23,031

REFERENCE/DOCKET NUMBER: GH30358

TELECOMMUNICATION INFORMATION:

TELEPHONE: 610-407-0700

TELEFAX: 610-407-0701

TELEX: 846169

INFORMATION FOR SEQ. ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1109 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: cDNA

US-09-088-651-1

alignment_scores:

Quality: 59.00 Length: 11

Ratio: 6.556 Gaps: 0

Percent Similarity: 81.818 Percent Identity: 63.636

alignment_block:

US-09-444-281-36 x US-09-088-651-1 ..

Align seg 1/1 to: US-09-088-651-1 from: 1 to: 1109

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seq_name: /cgn2_6/ptodata/2/ina/6B_COMB.seq:US-09-088-651-6

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APPLICANT: SAMPATH, Kuber T.
TITLE OF INVENTION: Morphogenic Protein-Specific Cell
TITLE OF INVENTION: Surface Receptors and Uses Therefor
NUMBER OF SEQUENCES: 18
CORRESPONDENCE ADDRESS:
ADDRESSEE: Testa, Hurwitz & Thibault
STREET: 125 High St.
CITY: Boston
STATE: MA
COUNTRY: USA
ZIP: 02110

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/481,337A
FILING DATE: 02-JUN-1995
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: MEYERS, Thomas C.
REGISTRATION NUMBER: 36,989
REFERENCE/DOCKET NUMBER: CRP-097CP2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 248-7000
TELEFAX: (617) 248-7100
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 1509 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
FEATURE:
NAME/KEY: CDS
LOCATION: 1..1509
OTHER INFORMATION: /product= "Human ALK1"
US-08-481-337A-1

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US-09-444-281-36 x US-08-481-337A-1 ..

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4 TrpProTrrTrpProTrr 9
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389 TGGCCCTGCTGCTGCTG 406

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Date: Jan 4, 2002 10:56 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

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-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0
-MAXLEN=2000000000 -USER=US09444281@CGN1.1_5145 -NCPU=6
-ICPU=3 -LONGLOG -NO_XLPHY -WAIT -THREADS=1

Search information block:

Query: US-09-444-281-36
Query length: 12
Database: GenEmbl.*
Database sequences: 1472140
Database length: 341344837
Search time (sec): 1728.450000

Score list:

Sequence	Strd Orig	ZScore	Escore	Len	Documentation
gb_com:BTINDLCD	+	70.00	149.45	4.77	550
gb_pat:AX098418	+	70.00	132.44	42.28	6446
gb_ro:MM08210	+	69.00	136.50	25.10	2651
gb_ro:AF289665	+	69.00	110.92	667.65	107257
gb_hlg:AP003754	-	69.00	109.64	786.64	129052
gb_hlg:AC091250	+	69.00	106.59	1.2e+03	200849
gb_ro:AF210429	+	67.00	144.52	8.98	456
gb_ro:AF166097	+	67.00	138.82	18.65	1040
gb_pl:ATPHTB	+	64.50	124.98	115.87	3850
gb_pl:ATPHTOCH	+	64.50	103.05	184.57	6509
gb_pl:AC005724	+	64.50	99.67	1.8e+03	86671
gb_pl:AP003141	+	64.50	103.05	1.8e+03	141275
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gb_hlg:AC027113	+	64.00	97.80	3.6e+03	159391
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gb_ro:RNYJ7153	+	63.00	118.97	237.92	5523
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gb_ba:AE004649	+	63.00	108.98	856.61	23425
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seq_documentation_block:

LOCUS BTINDLCD 550 bp mRNA MAM 07-OCT-1992
DEFINITION B.taurus mRNA for indolicidin.
ACCESSION X67340
VERSION X67340.1 GI:462
KEYWORDS indolicidin.
SOURCE cow.
ORGANISM Bos taurus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
Bovidae; Bovinae; Bos.

REFERENCE

1 (bases 1 to 550)
Del Sal, G., Storici, P., Schneider, C., Romeo, D. and Zanetti, M.
Bovidae. Biochem. Biophys. Res. Commun. 187 (1), 467-472 (1992)

MEDLINE

2 (bases 1 to 550)
Del Sal, G.
Direct Submission
Submitted (20-JUL-1992) G. Del Sal, Univ. of Trieste, Dip. di
Biochimica, Biofisica e Chimica, delle Macromolecole, Via A.
Valerio, 38, 34127 Trieste, ITALY

FEATURES

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Percent Similarity: 100.000 Percent Identity: 88.889

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DEFINITION Sequence 5 from Patent WO0119969.
ACCESSION AX098418
VERSION AX098418.1 GI:13537710
KEYWORDS Nicotiana Benthamiana.

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ORGANISM      Nicotiana benthamiana
               Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
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               Asteridae; euasterids I; Solanales; Solanaceae; Nicotiana.
REFERENCE      JOURNAL
AUTHORS
TITLE
FEATURES
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Percent Similarity: 100.000      Percent Identity: 88.889
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DEFINITION Mus musculus tropoelastin mRNA, complete cds.
ACCESSION  U08210
VERSION    U08210.1 GI:473273
KEYWORDS
SOURCE
ORGANISM
house mouse.
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 2651)
Wydner,K.S., Sechler,J.L., Boyd,C.D. and Passmore,H.C.
Use of an inltron polymorphism to localize the tropoelastin gene to
mouse chromosome 5 in a region of linkage conservation with human
chromosome 7
Genomics 23 (1), 125-131 (1994)
95130069
2 (bases 1 to 2651)
Boyd,C.D.
Direct Submission
Submitted (30-MAR-1994) Charles D. Boyd, Department of Surgery,
UMDNJ - Robert Wood Johnson Medical School, 51 French St., New
Brunswick, NJ 08903, USA
Location/Qualifiers
1. .2651
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3 ArgTTPProTTPTrProTparargTlys 12				
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seq_documentation_block:				
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DEFINITION	Mus musculus Eif4h gene, partial cds; LIMK1 gene, complete cds; and			14-AUG-2000
ACCESSION	AF289665			
VERSION	AF289665.1	GI:9800517		
KEYWORDS	house mouse.			
SOURCE	Mus musculus.			
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
REFERENCE	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.			
TITLE	1 (bases 1 to 107257)			
AUTHORS	Green, E.D.			
JOURNAL	Direct Submission			
	Submitted (26-JUL-2000) Genome Technology Branch, National Human			
	Genome Research Institute, 49 Convent Dr. Rm. 2A02, Bethesda, MD			
	20892, USA			
FEATURES	Location/Qualifiers			
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seq_documentation_block:
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DEFINITION Mus musculus chromosome 5 clone RP23-315E2 strain C57BL6/J, WORKING
DRAFT SEQUENCE, 7 unordered pieces.
AC091250
AC091250.1 GI:13592171
HTG: HTGS_PHASE1, HTGS_DRAFT.
KEYWORDS house mouse.
SOURCE
ORGANISM

REFERENCE
AUTHORS
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 200849)

Ayala, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W.,
Bouffard, G.G., Brinkley, C., Brooks, S., Dietrich, N.L., Granite, S.,
Guan, X., Gupta, J., Ho, S.-L., Idol, J.R., Karlins, E., Lee-Lin, S.-Q.,
Legaspi, R., Lim, M., Maduro, Q.L., Maduro, Y.B., Masello, C.,
Mastrian, S.D., McCloskey, J.C., McDowell, J., Pearson, R., Prasad, A.,
Shaychenko, Y., Snyder, B., Stantrilpop, S., Thomas, J.W., Thomas, P.D.,
Tingstrom, E.E., Touchman, J.W., Tsugeon, C., Vogt, J.L., Walker, M.A.,
Wetherby, K.D., Zhang, L.-H. and Green, E.D.

NIH Comparative Sequencing Initiative
Unpublished
2 (bases 1 to 200849)

Green, E.D.
Direct Submission
Submitted (11-APR-2001) NIH Intramural Sequencing Center, 8717
Groveomont Circle, Gaithersburg, MD 20877, USA

Genome Center
Center: NIH Intramural Sequencing Center

Center code: NISC
Web site: http://www.nisc.nih.gov

Contact: nisc-mouse@nih.gov
Project Information

Center project name: aty
Center clone name: 315E02

Summary Statistics

Sequencing vector: plasmid, n/a; 100% of reads
Chemistry: Dye-terminator Big Dye, 100% of reads

Assembly program: Phrap; version 0.990319
Consensus quality: 197727 bases at least Q40

Consensus quality: 198423 bases at least Q30
Consensus quality: 198823 bases at least Q20

Insert size: 20200; agarose-ftp
Insert size: 200249; sum-of-ctrls

Quality coverage: 9.87x in Q20 bases; agarose-ftp
Quality coverage: 9.95x in Q20 bases; sum-of-ctrls

* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 2812: contig of 2812 bp in length
2813 2912: gap of unknown length
2913 9301: contig of 6389 bp in length
9302 9401: gap of unknown length
9402 17173: contig of 7772 bp in length
17174 17273: gap of unknown length
17274 26044: contig of 8771 bp in length
26045 26144: gap of unknown length

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/strain="C57BL6/J"
/db_xref="taxon:10090"
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/clone_11b="RPCI mouse BAC library 23"
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26145..65929
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66030..109036
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BASE COUNT 53444 a 49164 c 49019 g 48614 t 608 others
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Quality: 69.00 Length: 10
Ratio: 7.667 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x AC091250 ..

Align seg 1/1 to: AC091250 from: 1 to: 200849

3 ArgTrpProtRTPProtRTPArgArgLys 12

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148453 AGTGGCCTTGTCGCTGACGCTCGG 148482

seq_name: gb_ro:AF210429

seq_documentation_block:
LOCUS AF210429 456 bp mRNA ROD 30-APR-2001

DEFINITION Mus musculus group X secretory phospholipase A2 (Plazg10) mRNA,
complete cds.

ACCESSION AF210429
VERSION AF210429.1 GI:12003290

KEYWORDS
SOURCE

ORGANISM

Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 456)

Morikawa, Y., Saiga, A., Yokota, Y., Suzuki, N., Ikeda, M., Ono, T.,
Nakano, K., Fujii, N., Ishizaki, J., Arita, H. and Hanasaki, K.

Mouse group X secretory phospholipase A2 induces a potent release
of arachidonic acid from spleen cells and acts as a ligand for the
phospholipase A2 receptor

Arch Biochem Biophys. 381 (1), 31-42 (2000)

JOURNAL
MEDLINE
PUBMED
20470496
11019817

REFERENCE
2 (bases 1 to 456)
Morikawa, Y., Saiga, A., Yokota, Y., Suzuki, N., Ikeda, M., Ono, T.,

AUTHORS

TITLE
JOURNALML

Nakano,K., Fujii,N., Ishizaki,J., Arita,H. and Hanasaki,K.
Direct Submission
Submitted (02-DEC-1999) Shionogi Research Lab, Fukushima-ku Sagisu
5-12-4, Osaka 553-0002, Japan

FEATURES
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Location/Qualifiers
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/db_xref="taxon:10090"

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/db_xref="GI:12003291"
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BASE COUNT
ORIGIN
98 a 125 c 129 g 104 t

alignment_scores:
Quality: 67.00 Length: 10
Ratio: 7.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-09-444-281-36 x AF210429
Align seg 1/1 to: AF210429 from: 1 to: 456

seq_name: gb_ro:AF166097

seq_documentation_block:
LOCUS AF166097 1040 bp mRNA ROD 06-DEC-1999
DEFINITION Mus musculus group X secreted phospholipase A2 (Pla2g10) mRNA,
complete cds.

ACCESSION AF166097
VERSION AF166097.2 GI:6525307

KEYWORDS house mouse;
SOURCE Mus musculus
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE AUTHORS Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
On the diversity of secreted phospholipases A(2). Cloning, tissue distribution, and functional expression of two novel mouse group III enzymes
J. Biol. Chem. 274 (44), 31195-31202 (1999)

JOURNAL MEDLINE Erratum:[published erratum appears in J Biol Chem 2000 Jan 21;275(3):22461]
REMARK 2 (bases 1 to 1040)
Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
Direct Submission
Submitted (07-JUL-1999) IPMC, CNRS, 660, route des Lucioles, Valbonne 06560, France
3 (bases 1 to 1040)
Valentin,E., Ghomashchi,F., Gelb,M.H., Lazdunski,M. and Lambeau,G.
Direct Submission
Submitted (06-DEC-1999) IPMC, CNRS, 660, route des Lucioles, Valbonne 06560, France
Sequence update by submitter

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COMMENT      On Dec 6, 1999 this sequence version replaced gi:6164695.
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        /CGAENKQCLLCRCDELAACLAGFTYHLKYLFPSPILCEKSPKCN"
BASE COUNT   244 a      258 c      277 g      261 t
ORIGIN
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  Ratio:      7.444      Gaps:    0
Percent Similarity: 90.000      Percent Identity: 80.000
alignment_block:
US-09-444-281-36 x AF166097 ..
Align seg 1/1 to: AF166097 from: 1 to: 1040
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      330 CTGTTATGTGGCCTTGTCGCATGAGAGA 359
seq_name: gb_pl:ATPHYB
seq_documentation_block:
LOCUS       ATPHYB      3850 bp      mRNA      PLN      13-SEP-1994
DEFINITION  Arabidopsis thaliana phyB mRNA for phytochrome.
ACCESSION   X17342
VERSION     X17342.1 GI:16422
KEYWORDS    phytochrome; phytochrome.
SOURCE      thale cress.
ORGANISM    Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
1 (bases 1 to 3850)
Sharrock,R.
Direct Submission
Submitted (21-DEC-1989) Sharrock, R., Montana State University,
Department of Biology, Bozeman, MT 59717 USA
2 (bases 1 to 3850)
Sharrock,R.A. and Quail,P.H.
Novel phytochrome sequences in Arabidopsis thaliana: structure,
evolution, and differential expression of a plant regulatory
photoreceptor family
Genes Dev. 3 (11), 1745-1757 (1989)
90108670
Location/Qualifiers
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    /strain="Columbia"
    /db_xref="taxon:3702"
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  93..3611
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ORIGIN

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Quality: 64.50 Length: 11
Ratio: 6.450 Gaps: 1
Percent Similarity: 90.909 Percent Identity: 81.818

alignment_block:
US-09-444-281-36 x ATHPTOCHB ..
Align seg 1/1 to: ATHPTOCHB from: 1 to: 6509

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2325 CGGTGGCCGTGGCGGTGGCGGTGGCGAGAGA 2357

seq_name: gb_pl::AC005724

seq_documentation_block:
LOCUS AC005724 86671 bp DNA PLN 05-APR-2000
DEFINITION Arabidopsis thaliana chromosome II section 109 of the complete sequence. Sequence from clones MSF3.
ACCESSION AC005724 AE002093
VERSION AC005724.2 GI:6598484
KEYWORDS HTG.
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
REFERENCE Arabidopsis thaliana
AUTHORS Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

1 (bases 1 to 86671)
Lin,X., Kaul,S., Rounsley,S.D., Shea,T.P., Benito,M.-I., Town,C.D., Fujii,C.Y., Mason,T.M., Bowman,C.L., Barnstead,M.E., Feldhuth,T.V., Buell,C.R., Ketchum,K.A., Lee,J.J., Ronning,C.M., Ko,H., Moffatt,K.S., Cronin,L.A., Shen,M., VanAken,S.E., Umayam,L., Patton,L.J., Gill,J.E., Adams,M.D., Carrera,A.J., Creasy,T.H., Goodman,H.M., Somerville,C.R., Copenhagen,G.P., Preuss,D., Nierman,W.C., White,O., Eisen,J.A., Salzberg,S.L., Fraser,C.M. and Venter,J.C.
TITLE Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana
JOURNAL Nature 402 (6763), 761-768 (1999)
MEDLINE 20083487
PubMed 10617197
REFERENCES 2 (bases 1 to 86671)
Lin,X.
DIRECT SUBMISSION Direct Submission
SUBMITTED (09-MAR-2000) The Institute for Genomic Research, 9712 Medical Center Dr., Rockville, MD 20850, USA
ON DEC 17, 1999 this sequence version replaced gi:4185128.
The sequence and annotation of chromosome 2 were merged from those of the individual clones on this chromosome after removing overlaps. For detailed information, please see the TIGR web site (<http://www.tigr.org/tdb/at/cat.html>).
COMMENT GENES were identified by a combination of three methods: Gene prediction programs including GRAIL (<ftp://arthur.epm.ornl.gov/pub/grail/>), GeneFinder (Phil Green, University of Washington), GENSCAN (Chris Burge, <http://genome.stanford.edu/EBNCAM.html>), and NetPlantGene (<http://www.cbs.dtu.dk/services/NetGene2/>), searches of the complete sequence against a peptide database and plant EST databases at TIGR, and manual curations based on those analyses.

Annotated genes are named to indicate the level of evidence for their annotation. Genes with similarity to other proteins are named after the database hits. Genes without significant peptide similarity but with EST similarity are named as 'unknown' proteins. Genes without protein or EST similarity, that are predicted by two or more gene prediction programs over most of their length are annotated as 'hypothetical' proteins. Genes encoding tRNAs are predicted by tRNAscan-SE (Sean Eddy, <http://genome.wustl.edu/eddy/tRNAscan-se/>). Simple repeats were identified by repeatmasker (Arrian Smil, <http://ftp.genome.washington.edu/RM/RepeatMasker.html>). Genes are numbered from the top to bottom of the chromosome.

We thank the CSHL/WashU/ABI consortium for sequencing BAC clones F6p23, F5j6, T17A5, and T13L16, the ESSA group for sequencing clone F13D4, and Scott Jackson, Jiming Jiang, Klaus Meyer, Eric Richards and Satoshi Tabata for helpful assistance. In addition, we would like to thank the TIGR Bioinformatics Department, especially Lixin Zhou, Hanif Khalak, Michael E. Heaney, Lily Fu, Feng Liang, Jeremy Peterson, Michael Holmes, and Delwood Richardson for software and database support.

This work was supported by the National Science Foundation, Department of Energy and the US Department of Agriculture.

Address all correspondence to: at@tigr.org.

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FEATURES
source
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    /organism="Arabidopsis thaliana"
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    RKYKTELELKRFPKVPKTEFFPALDLYVKQVMMLEHLKRLDKRLRNKIKT
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    DLPGMNDLDRGKPTTHKVFSGEVAISGALLALAFEHLEADVSCKMVRVKEEL
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    HSPGLCRSNLSGSESHNNSSSYLVLESGSRBMVPVLESNOLGADVNDSDS
    TRISGRSCDPCDMDGLDEKYEVLKVKLFPNIDHVGSGPOKKNISJGNSKND
    GRCLSMQSTYIYIDQENTLKVHYSTKLSGKDRVPSPHRYMSECGDPYKGIKESV
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           /label=ex14
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           /label=ex17
           /number=17
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           /label=ex18
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           /label=ex19
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alignment_scores:
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  ratio: 8.000      gaps: 0
  percent similarity: 88.889      percent identity: 77.778

alignment block:
US-09-444-281-36 x MMA2IXCOA/rev ..
Align seg 1/1 to reverse of: MMA2IXCOA from: 1 to: 19479
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4 TrpProTrrPrprrPrprrArgArgLys 12
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4355 TGCCCTGTGGTGGCTGGAGACCGG 4329

seq_name: gb_hctg:AL356097_0

seq_documentation_block:

WPCOMMENT

Sequence split into 4 fragments LOCUS AL356097 Accession AL356097

Fragment Name Begin End

AL356097_0 1 110000

AL356097_1 100001 210000

AL356097_2 200001 310000

AL356097_3 300001 355026

LOCUS AL356097 355026 bp DNA HTG 07-APR-2001

DEFINITION Homo sapiens chromosome 1 clone RP11-180A14, *** SEQUENCING IN

PROGRESS ***, 38 unordered pieces.

ACCESSION AL356097 AC058797

VERSION AL356097.11 GI:11414557

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 355026)

Plumb,B.

Direct Submission

Submitted (07-APR-2001) Sanger Centre, Hinxton, Cambridgeshire,

CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On May 15, 2001 this sequence version replaced gi:8077023

gi:9797422.

----- Genome Center

Center: Sanger Centre

Center code: SC

Web site: http://www.sanger.ac.uk

Contact: humquerry@sanger.ac.uk

----- Project Information

Center project name: bA180A14

----- Summary Statistics

Assembly program: XGAP4; version 4.5

Sequencing vector: M13; M7815; 43% of reads

Sequencing vector: plasmid; L08752; 56% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Consensus quality: 33302 bases at least Q40

Consensus quality: 346727 bases at least Q20

Insert size: 351326; sum-of-contigs

Insert size: 176896; 2.8% error; agarose-fp

Quality coverage: 3.95x in Q20 bases; sum-of-contigs Quality

coverage: 8.06x in Q20 bases; agarose-fp

Draft Sequence Produced by Whitehead Institute/MIT Center for

Genome Research, 320 Charles Street,

Cambridge, MA 02141, USA.

http://www-seq.wi.mit.edu.

* NOTE: This is a 'working draft' sequence. It currently

* consists of 38 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

1 16446: contig of 16446 bp in length

16447 16546: gap of 100 bp

16547 22746: contig of 6200 bp in length

22747 22846: gap of 100 bp

22847 45482: contig of 22636 bp in length

45483 45582: gap of 100 bp

45583 49251: contig of 3665 bp in length

49252 49351: gap of 100 bp

49352 53339: contig of 3888 bp in length

FEATURES

Source

* 53240 53339: gap of 100 bp

* 53340 58178: contig of 4839 bp in length

* 58179 58278: gap of 100 bp

* 58279 71754: contig of 13476 bp in length

* 71755 71854: gap of 100 bp

* 71855 76641: contig of 4787 bp in length

* 76642 76741: gap of 100 bp

* 76742 80415: contig of 3674 bp in length

* 80416 80515: gap of 100 bp

* 80516 89902: contig of 9387 bp in length

* 89903 90002: gap of 100 bp

* 90003 93661: contig of 3659 bp in length

* 93662 93761: gap of 100 bp

* 93762 101785: contig of 8024 bp in length

* 101786 101885: gap of 100 bp

* 101886 109333: contig of 7448 bp in length

* 109334 109433: gap of 100 bp

* 109434 121288: contig of 11855 bp in length

* 121289 121388: gap of 100 bp

* 121389 128767: contig of 7379 bp in length

* 128768 128867: gap of 100 bp

* 128868 131917: contig of 3050 bp in length

* 131918 132017: gap of 100 bp

* 132018 138795: contig of 6778 bp in length

* 138796 138895: gap of 100 bp

* 138896 144660: contig of 5765 bp in length

* 144661 144760: gap of 100 bp

* 144761 156587: contig of 11827 bp in length

* 156588 156687: gap of 100 bp

* 156688 158939: contig of 2252 bp in length

* 158940 159039: gap of 100 bp

* 159040 161353: contig of 2314 bp in length

* 161354 161453: gap of 100 bp

* 161454 165170: contig of 3717 bp in length

* 165171 165270: gap of 100 bp

* 165271 168479: contig of 3209 bp in length

* 168480 168579: gap of 100 bp

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* 172189 172288: gap of 100 bp

* 172289 177084: contig of 4796 bp in length

* 177085 177184: gap of 100 bp

* 177185 199876: contig of 22692 bp in length

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* 240236 240335: gap of 100 bp

* 240336 263715: contig of 23380 bp in length

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* 307329 307428: gap of 100 bp

* 307429 313985: contig of 6557 bp in length

* 313986 314085: gap of 100 bp

* 314086 330085: contig of 16000 bp in length

* 330086 330185: gap of 100 bp

* 330186 334887: contig of 4702 bp in length

* 334888 334987: gap of 100 bp

* 334988 355026: contig of 20039 bp in length.

Location/Qualifiers

1.355026

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Ratio: 8.000	9
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Align seg 1/1 to: AP001366 from: 1 to: 146081

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Mon, Jan 7 10:42:19 2002

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us-09-444-281-36.rge

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